

GenCore version 5.1.6
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OM protein - nucleic search, using frame_plus_p2n model

Run on: November 7, 2004, 04:47:28 ; Search time 4615 Seconds

(without alignments)
2090.379 Million cell updates/sec

Title: US-10-071-174-2

Perfect score: 1084

Sequence: 1 MVDQLERTTADPLRERTE.....QAFLSCLLTAFIYLWTELL 204

Scoring table: BLOSUM62

Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 4526729 seqs, 23644849745 residues

Total number of hits satisfying chosen parameters: 9053458

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:

-MODEL=frame+ p2n.model -DEV=xlp
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-DB=GenEmbl -QFMT=fastp -SUFFIX=rg -MINMATCH=0.1 -LOOFC=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=blosum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -THR SCORE=pct -THR MAX=100 -THR MIN=0 -ALIGN=15 -MODE=LOCAL
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-DSV TIMEOUT=120 -WARN TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

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2: gb_htg:*
3: gb_in:*
4: gb_cm:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pli:*
9: gb_pr:*
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11: gb_scs:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	1084	100.0	615	9 AF285092	AF285092 Homo sapi
2	1084	100.0	887	9 AF326964	AF326964 Homo sapi
3	1045	96.4	1168	6 BD233466	BD233466 Human pro
4	1036.5	95.6	726	6 CQ752105	CQ752105 Sequence

5	1035	95.5	582	6 BD233456	BD233456 Human pro
6	1035	95.5	585	9 HSA458330	HSA458330 Homo sapi
c 7	855	78.9	93287	2 AC023906	AC023906 Homo sapi
8	494	45.6	214669	2 AC018903	AC018903 Homo sapi
9	487	44.9	1074	10 AY029163	AY029163 Rattus no
10	474.5	43.8	1209	10 AF102501	AF102501 Mus muscu
11	474.5	43.8	1225	10 AF067660	AF067660 Mus muscu
12	467.5	43.1	1257	10 BC052690	BC052690 Mus muscu
13	342	31.5	240461	2 AC111669	AC111669 Rattus no
14	333.5	30.8	169914	2 AC133947	AC133947 Mus muscu
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17	193.5	17.9	1475	5 CCNR13	CCNR13 C. corunxi
18	188.5	17.4	534	5 AF375661	AF375661 Gallus ga
19	179	16.5	531	5 AF441285	AF441285 Danio rer
c 20	178.5	16.5	151102	14 AF282130	AF282130 Meleagrid
21	178.5	16.5	159160	14 AF291866	AF291866 Meleagrid
c 22	178.5	16.5	159160	14 AF291866	AF291866 Meleagrid
c 23	144	13.3	170193	5 AL935189	AL935189 Zebrafish
24	143.5	13.2	636	5 AF120210	AF120210 Gallus ga
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28	131	12.1	694	4 AB080230	AB080230 Canis fam
29	129	11.9	658	6 AX525910	AX525910 Sequence
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37	127	11.7	791	6 BD203742	BD203742 Human nuc
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41	126	11.6	646	4 AB080724	AB080724 Felis cat
42	126	11.6	677	9 AF007826	AF007826 Homo sapi
43	125	11.5	525	5 AF120211	AF120211 Gallus ga
44	125	11.5	624	6 AR027309	AR027309 Sequence
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ALIGNMENTS

RESULT 1	AF285092	Homo sapiens Bcl-2-like protein 10 mRNA, complete cds.	615 bp	mrna	linear	PRI 08-NOV-2001
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DEFINITION	AF285092					
ACCESSION	AF285092					
VERSION	AF285092.1	GI:9837265				
KEYWORDS						
SOURCE						
ORGANISM						
REFERENCE						
AUTHORS						
TITLE						
JOURNAL						
MEDLINE						
PUBMED						
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AUTHORS						
TITLE						
JOURNAL						
FEATURES						
source						


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Db      350 GTGACCTTCGCGAGGAGCGTGTGTGAGAGAGGGCGCTGGTGACCGCCGGTGGGAAG 409
Qy      121 TrpGlyPheGlnProArgLeuLysGluGlnGluGlyAspValAlaArgAspCysGlnArg 140
Db      410 TGGGGCTTCAGCGCGGCTAAAGGAGCAGGAGGGCGAGCTGCCCGGGAGCTGCCAGGCG 469
Qy      141 LeuValAlaLeuLeuSerArgLeuMetGlyGlnHisArgAlaTrpLeuGlnAlaGln 160
Db      470 CTGGTGGCTTCTGAGCTCGCGGCTCAATGGGGCAGCACCGCGCTGCTGCGAGCTCAG 529
Qy      161 GlyGlyTrpAspGlyPheCysHisPhePheArgTrpProPheProLeuAlaPheTrpArg 180
Db      530 GCGGGCTGGATGGCTTTGTCTACCTCTTCAGAGACCCCTTCCACTGGCTTTTGGAGA 589
Qy      181 LysGlnLeuValGlnAlaPheLeuSerCysLeuLeuThrThrAlaPheLeuTrpLeuTrp 200
Db      590 AAACAGCTGGTCCAGGCTTTTCTGTCTGTCTGTCTGTCTGTCTGTCTGTCTGTCTGTCT 649
Qy      201 ThrArgLeuLeu 204
Db      650 ACACGATTATTA 661

RESULT 3
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LOCUS      Human protein having hydrophobic domain and DNA encoding the same.
DEFINITION
ACCESSION BD233466
VERSION    BD233466.1 GI:33043236
KEYWORDS  JP 2002519016-A/12.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
REFERENCE  Kato,S. and Kimura,T.
AUTHORS   Mammalia; Eutheria; Primates; Catarrhini, Hominidae; Homo.
TITLE     1 (bases 1 to 1168)
JOURNAL   Human protein having hydrophobic domain and DNA encoding the same
COMMENT   Patent: JP 2002519016-A 12 02-JUL-2002;
          SAGAMI CHEMICAL RESEARCH CENTER, PROTEGENE INC
          OS Homo sapiens (human)
          PN JP 2002519016-A/12
          PD 02-JUL-2002 JP 2000557267
          PF 18-JUN-1999 JP 2000557267
          PI SEISHI KATO,TOMOKO KIMURA
          PC

C12N15/09, C07K14/47, C12N1/15, C12N1/19, C12N5/10, C12N5/00, C12N5/ PC
CC Human protein having hydrophobic domain and DNA encoding the
CC same
FH Key same Location/Qualifiers
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Best Local Similarity: 100.00% Mismatches: 0
Query Match: 96.40% Indels: 0
DB: 6 Gaps: 0
US-10-071-174-2 (1-204) x BD233466 (1-1168)

Qy      9 ThrThrMetAlaAspProLeuArgGluArgThrGluLeuLeuAlaAspTyrLeuGly 28
Db      1 ACCACATGGCGACCGCTGGGGAGCGCACCGAGCTGTTCGCGGACCTACCTGGGG 60
Qy      29 TyrCysAlaArgLupProGlyThrProGluProAlaProSerThrProGluAlaVal 48
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Db      121 CTGGCTCCCGGGCGCGCAGTTACGGCAGATTACCGGTCCTTTTCTCCGCTACCTC 180
Qy      69 GlyTyrProGlyAsnArgPheGluLeuValAlaLeuValAlaAspSerValLeuSerAsp 88
Db      181 GGCTACCCCGGGAAACCGCTTCGAGCTGGTGCGCTGATGGCGGATTCCGTGCTCTCCGAC 240
Qy      89 SerProGlyProThrTrpGlyArgValValThrLeuValThrPheAlaGlyThrLeuLeu 108
Db      241 AGCCCCGGCCCCACCTGGGGCAGAGTGGTGACCTCTGTGACCTTCGACGAGGAGCTGCTG 300
Qy      109 GluArgGlyProLeuValThrAlaArgTrpLysTrpGlyPheGlnProArgLeuLys 128
Db      301 GAGAGAGGGCGGCTGGTGACCGCGCGGTGAAGAAGTGGGGCTTCAGCGCGCGCTAAAG 360
Qy      129 GluGlnGluGlyAspValAlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArg 148
Db      361 GAGCAGAGGGCGGACGTCGCCCGGGACTGCCAGCGCTGGTGCGCTTGTGAGCTGCGGG 420
Qy      149 LeuMetGlyGlnHisArgAlaTrpLeuGlnAlaGlnGlyTrpAspGlyPheCysHis 168
Db      421 CTCATGGGGCAGCACCGCGCTGGCTGACGCTCAGGGCGCTGGGATGGCTTTTGTAC 480
Qy      169 PhePheArgThrProPheProLeuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeu 188
Db      481 TTCTTCAGSACCCCTTTCACCTGGCTTTTGGAGAAACAGCTGGTCCAGGCTTTTCTG 540
Qy      189 SerCysLeuLeuThrThrAlaPheLeuTyrLeuTrpThrArgLeuLeu 204
Db      541 TCATGCTTGTAAACACAGCGCTTCATTATCTCTGGACAGATTATTA 588

RESULT 4
CQ752105
LOCUS      Sequence 38039 from Patent WO02068579.
DEFINITION
ACCESSION CQ752105
VERSION    CQ752105.1 GI:42387450
KEYWORDS
SOURCE     Homo sapiens (human)
ORGANISM  Homo sapiens
REFERENCE  Venter,C.J., Adams,M.C., Li,P.W. and Myers,E.W.
AUTHORS   Kits, such as nucleic acid arrays, comprising a majority of
TITLE     humanexons or transcripts, for detecting expression and other uses
          thereof
JOURNAL   Patent: WO 02068579-A 38039 06-SEP-2002;
          PE Corporation (NY) (US)
FEATURES
source 1..726
          Location/Qualifiers
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          /mol_type='unassigned DNA'
          /db_xref='taxon:9606'
ORIGIN
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Pred. No.: 4.66e-77 Length: 726
Score: 1036.50 Matches: 203
Percent Similarity: 89.04% Conservatives: 0
Best Local Similarity: 89.04% Mismatches: 1
Query Match: 95.62% Indels: 24
DB: 6 Gaps: 1
US-10-071-174-2 (1-204) x CQ752105 (1-726)

Qy      1 MetValAspGlnLeuArgGluArgThrThrMetAlaAspProLeuArgGluThrGlu 20
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QY 21 LeuLeuLeuAlaAspTyrLeuGlyTyrCysAlaArgGluProGlyThrProGluProAla 40
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QY 41 ProSerThrProGluAlaValLeuArgSerAlaAlaAlaArgGlnIleHis 60
DB 121 CCATCCACGCCCGAGCGCGCTGCGCTCGCGCGCGCGCGAGGTACCGGAGATTAC 180
QY 61 ArgSerPhePheSerAlaTyrLeuGlyTyrProGlyAsnArgPheGluLeuValAlaLeu 80
DB 181 CGGTCTTTTCTCCGCTACCTCGCTACCGCGCGGAACCGCTTCGAGCTGGCGCTG 240
QY 81 MetAlaAspSerValLeuSerAspSerProGlyProThrTyrGlyArgValThrLeu 100
DB 241 ATGGCGGATTCCGTCTCTCCGACAGCCCGCGCGCGCGCGCGCGCGCTGCGCTC 300
QY 101 ValThrPheAlaGlyThrLeuLeuGluArgGlyProLeuValThrAlaArgTyrIleHis 120
DB 301 GTGACCTTCGACGGGAGCTGCTGGAGAGAGGCGCGCTGGTGACCGCGCGTGGAGAG 360
QY 121 TrpGlyPheGlnProArgLeuLysGluGlnGluGlyAspValAlaAlaArgAspCysGlnArg 140
DB 361 TGGGGCTTCAGCGCGCGGTAAAGAGGACAGAGGCGCGCGCTGCGCGCGGACTGCCAGCGC 420
QY 141 LeuValAlaLeuSerSerArgLeuMetGlyGlnHisArgAlaTyrLeuGlnAlaGln 160
DB 421 CTGGTGCCCTTGCTGAGCTCGCGCTCATGGGCGACACCGCGCTGGCTGGCGCTCAG 480
QY 161 GlyGlyTrp----- 163
DB 481 GCGGCTGGGTGAGCAGCGCGCGGACACCGGCGGACCGGCGGCGGCGGCGGAG 540
QY 164 -----AspGlyPheCysHisPheArgThrProPheProLeuAl 177
DB 541 CGGCCAGAGGCTGGCAGCGATGGCTTTTGTCACTTTCAGAGACCCCTTTCCACTGGC 600
QY 177 aPheTyrArgLysGlnLeuValGlnAlaPheLeuSerCysLeuLeuThrThrAlaPheIle 197
DB 601 TTTTGGAGAAACAGCTGGTCCAGGCTTTCTGTATGCTGTTTAAACAGCCTTCAT 660
QY 197 eTyrLeuTyrThrArgLeuLeu 204
DB 661 TTATCTCTGACACGATTATTA 682

RESULT 5
BD233456 582 bp DNA linear PAT 17-JUL-2003
LOCUS Human protein having hydrophobic domain and DNA encoding the same.
DEFINITION BD233456
ACCESSION BD233456.1 GI:33043226
VERSION JP 2002519016-A/2.
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 582)
Kato, S. and Kimura, T.
Human protein having hydrophobic domain and DNA encoding the same
Patent: JP 2002519016-A 2 02-JUL-2002;
SAGAMI CHEMICAL RESEARCH CENTER, PROTEGENE INC
OS Homo sapiens (human)
PN JP 2002519016-A/2
PD 02-JUL-2002
PF 18-JUN-1999 JP 2000557267
PI SEISHI KATO, TOMOKO KIMURA
PC
C12N15/09, C07K14/47, C12N1/15, C12N1/19, C12N5/10, C12N5/00, C12N5/ PC
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CC Human protein having hydrophobic domain and DNA encoding the
CC same
PH Key Location/Qualifiers
FT source 1.582

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Location/Qualifiers
1.582
/organism='Homo sapiens'
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Pred. No.: 1035.00 Matches: 194
Score: 1035.00 Conservatives: 0
Percent Similarity: 100.00% Mismatches: 0
Best Local Similarity: 100.00% Indels: 0
Query Match: 95.48% Caps: 0
DB: 6
US-10-071-174-2 (1-204) x BD233456 (1-582)
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DB 1 ATGCCCGACCCGCTGCGGAGCGCACCGAGCTGTGCTGCGCGCTACCTGGGCTACTGC 60
QY 31 AlaArgGluProGlyThrProGluProAlaProSerThrProGluAlaAlaValLeuArg 50
DB 61 GCGCGGAACCGGACCCCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCGCG 120
QY 51 SerAlaAlaAlaArgLeuArgGlnIleHisArgSerPhePheSerAlaTyrLeuGlyTyr 70
DB 121 TCCGCGCGCGCGAGTTACGCGAGATTACCGGCTCTTTTCTCCGCTACCTCGGCTAC 180
QY 71 ProGlyAsnArgPheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerPro 90
DB 181 CCGCGGAACCGCTTCGAGCTGGTGGCGCTGATGCGGAGATTCCTGCTCTCCGACGCCC 240
QY 91 GlyProThrTyrGlyArgValValThrLeuValThrPheAlaGlyThrLeuLeuGluArg 110
DB 241 GCGCGCGCGCGCGCGAGTGGTGACGCTCGTACCTTCGACGAGCGCTGCTGGAGAGA 300
QY 111 GlyProLeuValThrAlaArgTyrLysLysLysLysLysLysLysLysLysLysLys 130
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QY 131 GluGlyAspValAlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArgLeuMet 150
DB 361 GAGGCGACGCTCGCGCGGACTGCCAGCGCTGGTGGCGCTGCTGAGCTCGCGGCTCATG 420
QY 151 GlyGlnHisArgAlaTyrLeuGlnAlaGlnGlyLysLysLysLysLysLysLysLys 170
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QY 171 ArgThrProPheProLeuAlaPheTyrArgLysGlnLeuValGlnAlaPheLeuSerCys 190
DB 481 AGGACCCCTTTCACCTGGCTTTTGGAGAAACAGCTGGTCCAGGCTTTTCTGTATGC 540
QY 191 LeuLeuThrThrAlaPheIleTyrLeuTyrThrArgLeuLeu 204
DB 541 TTGTTAAACAGCGCTTCATTTATCTCTGGACAGATTATTA 582

RESULT 6
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LOCUS Homo sapiens NRH gene for anti-apoptotic protein.
DEFINITION HSA458330
ACCESSION AJ458330
VERSION AJ458330.1 GI:20338765
KEYWORDS anti-apoptotic protein; NRH gene.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1
AUTHORS Aouacheria, A., Arnaud, E., Venet, S., Lalle, P., Gouy, M., Rigal, D. and
Gillet, G.
TITLE Nrhl, a human homologue of Nr-13 associates with Bcl-Xs and is an

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Alignment Scores:

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Query Match: 45.57% Indels: 2
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US-10-071-174-2 (1-204) x AC018903 (1-214669)

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Qy 21 LeuLeuLeuAlaAspTyrLeuGlyTyr-CysAlaArgGluProGlyThrProGluProAl 40
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PLPLGFWRRLLIQAFLSGFFATAIFFWKRL"

ORIGIN
Alignment Scores:
Pred. No.: 3,51e-30 Length: 1209
Score: 474.50 Matches: 94
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Best Local Similarity: 48.21% Mismatches: 56
Query Match: 43.77% Indels: 13
DB: 10 Gaps: 3

US-10-071-174-2 (1-204) x AF102501 (1-1209)
QY 13 AspProLeuArgGluArgThrGluLeuLeuAlaAspTyrLeuGlyTyrCysAlaArg 32
Db 132 GACCCACTGCATGACGCACTAGACGGCTGCTGTGACTACATATCTTCGCGCAGG 191
QY 33 GluProGlyThrProGluProAlaProSerThrProGluAlaAlaValLeuArgSerAla 52
Db 192 GAGCGGACACCCAGAGCACCGCCACGCTGTGCGAGCGGCTTGCCTCGCTCTGTG 251
QY 53 AlaAlaArgLeuArgGlnIleHisArgSerPheSerAlaTyrLeuGlyTyrProGly 72
Db 252 ACTAGGCATGATCAGCAGGACGACCAAGTATTTTCTCTCTCTCGCAAGCGGGCG 311
QY 73 AsnArgPheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerProGlyPro 92
Db 312 AATCGCCTGGAGCTGTGTGAACAGATGGCAGATAGTTGCTCTCCAAAGACCAAGCTC 371
QY 93 ThrTrpGlyArgValValThrLeuValThrPheAlaGlyThrLeuLeuGluArgGlyPro 112
Db 372 AGCTGGAGCAACTGTGTGATGCTCTGCGGCTTCGCGGAGCGCTTATGAATCAAGCCCT 431
QY 113 LeuValThrAlaArgTrpLysLysTrpGlyPheGlnProArgLeuLysGluGlnGly 132
Db 432 TACATGGCTGTACAGCAGAGAGG-----GATCTGGG 464
QY 133 Asp-----ValAlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArgLeu 149
Db 465 AATCGGTGCATAGTACCGGACGCTGCTCATAGTGAACCTTCTGTATATATCTGCTC 524
QY 150 MetGly---GlnHisArgAlaTrpLeuGlnAlaGlnGlyTrpAspGlyPheCysHis 168
Db 525 ATGGGCGGTGGCAGCCGCGCAGCTGAGGCTCTCGCGGCTGGGATGGCTTTTGGCGG 584
QY 169 PhePheArgThrProPheProLeuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeu 188
Db 585 TTCTTCAAGATCTTTTACCGCTCGGCTTCTGGAGAAGATTCGTATTCAGGCTTTCTG 644
QY 189 SerCysLeuLeuThrThrAlaPheIleTyrLeuTrpThrArgLeu 203
Db 645 TCAGGCTCTTTTGCAACAGCCCATCTTTTATCTGGAACGTTA 689

RESULT 11
AF067660
LOCUS
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DEFINITION Mus musculus Bcl-2 homolog (Diva) mRNA, complete cds.
ACCESSION AF067660
VERSION AF067660.1 GI:3955265
KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 1225)
AUTHORS Inohara,N., Gourley,T.S., Carrio,R., Muniz,M., Merino,J.,
Garcia,I., Koseki,T., Hu,Y., Chen,S. and Nunez,G.
TITLE Diva, a Bcl-2 homologue that binds directly to Apaf-1 and induces
BH3-independent cell death
JOURNAL J. Biol. Chem. 273 (49), 32479-32486 (1998)
MEDLINE 99047617
PUBMED 9829980
REFERENCE 2 (bases 1 to 1225)
AUTHORS Inohara,N. and Nunez,G.
TITLE Direct Submission
JOURNAL Submitted (20-MAY-1998) Department of Pathology, University of
Michigan Medical School, 1500 E. Medical Dr., Ann Arbor, MI 48109,
USA
FEATURES
source Location/Qualifiers
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/mol_type="mRNA"
/strain="B6D2F1/J"
/db_xref="taxon:10090"
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143. .718
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/notes="Diva"
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Best Local Similarity: 48.21% Mismatches: 56
Query Match: 43.77% Indels: 13
DB: 10 Gaps: 3

US-10-071-174-2 (1-204) x AF067660 (1-1225)
QY 13 AspProLeuArgGluArgThrGluLeuLeuAlaAspTyrLeuGlyTyrCysAlaArg 32
Db 158 GACCCACTGCATGACGCACTAGACGGCTGCTGTGACTACATATCTTCGCGCAGG 217
QY 33 GluProGlyThrProGluProAlaProSerThrProGluAlaAlaValLeuArgSerAla 52
Db 218 GAGCGGACACCCAGAGCACCGCCACGCTGTGCGAGCGGCTTGCCTCGCTCTGTG 277
QY 53 AlaAlaArgLeuArgGlnIleHisArgSerPheSerAlaTyrLeuGlyTyrProGly 72
Db 278 ACTAGGCATGATCAGCAGGACGACCAAGTATTTTCTCTCTCTCGCAAGCGGGCG 337
QY 73 AsnArgPheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerProGlyPro 92
Db 338 AATCGCCTGGAGCTGTGTGAACAGATGGCAGATAAGTTGCTCTCCAAAGACCAAGCTC 397
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QY 93 ThrTTPGlyArgValThrLeuValThrPheAlaGlyThrLeuLeuGluArgGlyPro 112
Db 398 AGCTGGAGCAACTGGTGTATGCTTGGCCCTTCGGGGAGCCCTTATGAATCAAGGCCCT 457
QY 113 LeuValThrAlaArgTrpLysLysTTPGlyPheGlnProArgLeuLysGluGlnGly 132
Db 458 TACATGGCTGTCAAGCAGAGAGG-----GATCTGGGG 490
QY 133 Asp-----ValAlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArgLeu 149
Db 491 AATCGTGTATAGTACCGGAGACTGCTCTCATAGTGAACCTTCTGTATAATCTGCTC 550
QY 150 MetGly---GlnHisArgAlaTrpLeuGlnAlaGlnGlyGlyTTPAspGlyPheCysHis 168
Db 551 ATGGGGGTGTCGGCCCGCCAGGCTGGAGGCTCTCGGCGCTGGAGTGGCTTTTGGCGC 610
QY 169 PhePheArgThrProPheProLeuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeu 188
Db 611 TTCTTCAAGAACTCTTATCGCTCGCTTCTGGAGAAGATTGCTGATTGAGCTTTTCTG 670
QY 189 SerCysLeuLeuThrAlaPheLeuTyrLeuTTPThrArgLeu 203
Db 671 TCAGGCTTCTTGCACAGCCATCTTTTATCTGGAACGTTA 715
RESULT 12
LOCUS BC052690 1257 bp mRNA linear ROD 08-OCT-2003
DEFINITION Mus musculus Bcl2-like 10, mRNA (cDNA clone MGC:60542 IMAGE:30052580), complete cds.
ACCESSION BC052690
VERSION BC052690.1 GI:30851238
KEYWORDS MGC.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 1257)
AUTHORS Strausberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G., Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D., Altschul,S.F., Zeeberg,B., Buetow,K.H., Schaefer,C.F., Bhat,N.K., Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,P., Diatchenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L., Scapleton,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L., Scheetz,T.E., Brownstein,M.J., Usdin,T.B., Toshiyuki,S., Carninci,P., Prange,C., Raja,S.S., Loquellano,N.A., Peters,G.J., Abramson,R.D., Mullahy,S.J., Bosak,S.A., McEwan,P.J., McKernan,K.J., Malek,J.A., Gunaratne,P.H., Richards,S., Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Hulyk,S.W., Villalón,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A., Fahey,J., Heiton,E., Kettelman,M., Madan,A., Rodriguez,S., Sanchez,A., Whitting,M., Madan,A., Young,A.C., Shevchenko,Y., Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D., Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M., Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smallos,D.E., Schnerch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Generation and initial analysis of more than 15,000 full-length human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
22388257
REFERENCE 2 (bases 1 to 1257)
AUTHORS Strausberg,R.
JOURNAL Direct Submission
TITLE Submitted (16-MAY-2003) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA
REMARK NIH-MGC Project URL: <http://mgc.nci.nih.gov>
COMMENT Contact: MGC help desk
Email: cgaps-x@mail.nih.gov
Tissue Procurement: Minoru Ko
CDNA Library Preparation: Yulan Piao and Minoru Ko (National Institute on Aging, NIH: <http://lgeun.grc.nia.nih.gov/cDNA/>)

CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LNLN)
DNA Sequencing by: National Institutes of Health Intramural Sequencing Center (NISC), Gaithersburg, Maryland;
Web site: <http://www.nisc.nih.gov/>
Contact: nisc_mgc@nih.gov
Akhtar,N., Ayele,K., Beckstrom-Sternberg,S.M., Benjamin,B., Blakesley,R.W., Bouffard,G.G., Breen,K., Brinkley,C., Brooks,S., Dietrich,N.L., Granite,S., Guan,X., Gupta,J., Haghighi,P., Hansen,N., Ho,S.-L., Karlins,E., Kwong,P., Loric,P., Legaspi,R., Maduro,Q.L., Masello,C., Maskeri,B., Mastrian,S.D., McCloskey,J.C., McDowell,J., Pearson,R., Stantropop,S., Thomas,P.J., Touchman,J.W., Tourgeon,C., Vogt,J.L., Walker,M.A., Wetherby,K.D., Wiggins,L., Young,A., Zhang,L.-H. and Green,E.D.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LNLN at: <http://image.llnl.gov>
Series: IRAC Plate: 112 Row: h Column: 18
This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 7304926.

FEATURES
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gene

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/db_xref="LocusID:12049"
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/db_xref="CDD:smart00337"

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Best Local Similarity: 47.69% Mismatches: 57
Query Match: 43.13% Indels: 13
DB: 10 Gaps: 3

US-10-071-174-2 (1-204) x BC052690 (1-1257)

QY 13 AspProLeuArgGluArgThrGluLeuLeuAlaAspTyrLeuGlyTyrCysAlaArg 32
Db 160 GACCACTGATGACCGCACTAGACGGCTGCTGCTGACTACATATTTCTCGGCACGG 219
QY 33 GluProGlyThrProGluProAlaProSerThrProGluAlaAlaValLeuArgSerAla 52
Db 220 GAGCGGACACCCAGAGCCACCGCCCGCTGCTGTCGAGCGGCGCTTGTCTCTGTG 279

* as soon as it is available and the accession number will
* be preserved.

1 9792: contig of 9792 bp in length
9793 9892: gap of unknown length
9893 118024: contig of 108132 bp in length
118025 118124: gap of unknown length
118125 236959: contig of 118334 bp in length
236960 237059: gap of unknown length
237060 238398: contig of 1340 bp in length
238399 238499: gap of unknown length
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118125..119133
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ORIGIN

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Query Match: 31.55% Indels: 13
DB: 2 Gaps: 2

US-10-071-174-2 (1-204) x AC111669 (1-240461)

Qy 11 MetAlaSpProLeuArgGluAArgThrGluLeuLeuAlaAspTyrLeuGlyTyrCys 30
Db 21810 ATGGTGCACCGCTGCGAGTCCGACTAGACGCTGCTGACTACATATTTCTGC 21869
Qy 31 AlaArgGluProGlyThrProGluProAlaProSerThrProGluAla-AlaValLeuAr 50
Db 21870 GCAGGGCGCGACACCCCTGAGCCACTGCCACGCTCTGTGAGCGCGCTGCTGC 21929
Qy 50 gSerAlaAlaAlaArgLeuArgGlnIleHisArgSerPhePheSerAlaTyrLeuGlyTy 70
Db 21930 CTCTGTGACTAGTACATCCACAGGAGCACCAGGATCTTTTCACTCTCTCGGACTA 21989
Qy 70 rProGlyAsnArgPheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerPr 90
Db 21990 CCAGGGACACCGCTGAGCTGTGTGACAGATGGCGGATGCTCTCCATGACCA 22049
Qy 90 cGlyProThrTrpGlyArgValValThrLeuValThrPheAlaGlyThrLeuLeuGluAr 110
Db 22050 AGAGTTCAACTGGCGCGCTGTGTGATGCTCTGCGCTTCTGTTGGGACGCTAATGA 22109
Qy 110 gGlyProLeuValThrAlaArgTrpLysLysTrpGlyPheGlnProArgLeuLysGluGl 130
Db 22110 AGACAGACTGTTAAGCGGAGG-----AGGGATCA 22139
Qy 130 nGluGlyAsp-----ValAlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerAr 148
Db 22140 AAGAAACCGTCTCTACTGAGCGAGACTGCTATCTCATAGTAGCTTGTGTACAATCG 22199
Qy 148 gLeuMetClyGlnHisArgAlaTrpLeuGlnAlaGlnGlyGlyTrp 163
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RESULT 14

AC133947

LOCUS

DEFINITION

AC133947

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AC133947 169914 bp DNA linear HTG 10-JUL-2004
Mus musculus chromosome 9 clone RP24-18912, WORKING DRAFT SEQUENCE,
3 unordered pieces.
AC133947
AC133947.2 GI:50199129
HTG; HTGS PHASE1; HTGS_DRAFT; HTGS_ACTIVEPIN.
Mus musculus (house mouse)
Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 169914)
Wilson, R.K.
The sequence of Mus musculus clone
Unpublished
2 (bases 1 to 169914)
McPherson, J.D. and Waterston, R.H.
Direct Submission
Submitted (20-SEP-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
3 (bases 1 to 169914)
Wilson, R.K.
Direct Submission
Submitted (10-JUL-2004) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
On Jul 10, 2004 this sequence version replaced gi:23238076.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu
Contact: submissions@wustl.edu
----- Project Information -----
Center project name: M_BB0189102
----- Summary Statistics -----
Sequencing vector: M13; 0%
Sequencing method: plasmid; 100%
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 167792 bases at least Q40
Consensus quality: 168148 bases at least Q30
Consensus quality: 168291 bases at least Q20
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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21305..86045
/note="assembly_name:Contig14"
86146..169914
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FEATURES

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misc_feature
misc_feature

Center clone name: 365_N_15

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      8422..8449
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Query Match: 30.77% Indels: 13
DB: 10 Gaps: 3

US-10-071-174-2 (1-204) x AC115880 (1-202851)
QY 13 AspProLeuArgGluArgThrGluLeuLeuAlaAspTyrLeuGlyTyrCysAlaArg 32
Db 30352 GACCCACTGCATGAACGACTAGACGGCTGCTCTGACTACATATTCCTTCGCGACGG 30411
QY 33 GluProGlyThrProGluProAlaProSerThrProGluAlaAlaValLeuAlaGSerAla 52
Db 30412 GAGCGGACACCCGAGAGCCGCCCTCTGCTCGAGGGCGCTTCTGCTCTGCTG 30471
QY 53 AlaAlaArgLeuArgGlnIleHisArgSerPhePheSerAlaTyrLeuGlyTyrProGly 72
Db 30472 ACTAGGCAGATCCAGCAGGAGCACCAGAAATTTTTCCTCTCTCGAAAGCGGGGC 30531
QY 73 AsnArgPheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerProGlyPro 92
Db 30532 AATCGCTGGAGCTGGTGAACAGATGCGAGATAAGTTGCTCTCCAAGACCAAGACTTC 30591
QY 93 ThrTrpGlyArgValValThrLeuValThrPheAlaGlyThrLeuLeuGluArgGlyPro 112
Db 30592 AGCTGGAGCCAACTGCTGATGCTCTGCGGGGACGCTTATGAATCAAGGCCCT 30651
QY 113 LeuValThrAlaArgTrpLysLysTrpGlyPheGlnProArgLeuLysGluGlnGly 132
Db 30652 TACATGGCTGTCAAGCAGAGAGG-----GATCTGGGG 30684
QY 133 Asp-----ValAlaArgAspCysGlnArgLeuValAlaLeuLeuSerArgLeu 149
Db 30685 AATCGTGTCTAGTGACCCGAGACTGCTGCTCATAGTAACCTTCTGTATAATCTGCTC 30744
QY 150 MetGly---GlnHisArgAlaTrpLeuGlnAlaGlnGlyTrp 163
Db 30745 ATGGGGCGTCGGCACCGCCGAGCTGGAGGCTCTCGCGCGCTGG 30789

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Job time : 4754 secs

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GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 6, 2004, 21:41:06 ; Search time 6234 Seconds
(without alignments)
6728.579 Million cell updates/sec

Title: US-10-071-174-1
Perfect score: 887
Sequence: 1 cgggccaagaaccagcga.....ctcttccttgactgaagaa 887

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 4526729 seqs, 23644849745 residues

Total number of hits satisfying chosen parameters: 9053458

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : GenEmbl.*

- 1: gb.ba.*
- 2: gb.htg.*
- 3: gb.in.*
- 4: gb.om.*
- 5: gb.ov.*
- 6: gb.pat.*
- 7: gb.ph.*
- 8: gb.pl.*
- 9: gb.pr.*
- 10: gb.ro.*
- 11: gb.sts.*
- 12: gb.sy.*
- 13: gb.un.*
- 14: gb.vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB	ID	Description
1	887	100.0	887	9	AF326964	AF326964 Homo sapi
2	812.4	91.6	1168	6	BD232466	BD232466 Human pro
3	615	69.3	615	9	AF285092	AF285092 Homo sapi
4	585	66.0	585	9	AF285092	AF285092 Homo sapi
5	582	65.6	582	6	BD232456	BD232456 Human pro
6	574.4	64.8	726	6	CQ752105	CQ752105 Sequence
7	539.2	60.8	93287	9	AC023906	AC023906 Homo sapi
8	345.6	39.0	214669	2	AC018903	AC018903 Homo sapi
9	327.5	36.9	214669	2	AC018903	AC018903 Homo sapi
10	259.2	29.2	1289	10	AF102501	AF102501 Mus muscu
11	259.2	29.2	1289	10	AF102501	AF102501 Mus muscu
12	257.6	29.0	1257	10	BC052690	BC052690 Mus muscu
13	244.2	27.5	1074	10	AY029163	AY029163 Rattus no
14	147	16.6	169914	2	AC133947	AC133947 Mus muscu
15	147	16.6	202851	10	AC115880	AC115880 Mus muscu
16	141.8	16.0	240451	2	AC111659	AC111659 Rattus no
17	95.4	10.8	243686	2	AC036430	AC036430 Rattus no
18	85.2	9.6	160755	9	AC104976	AC104976 Homo sapi
19	68	7.7	2000	6	AX655393	AX655393 Sequence

C 20	62.2	7.0	295150	1	SC093126	AL939126 Streptomy
C 21	61.2	6.9	125020	9	AF429315	AF429315 Homo sapi
C 22	60	6.8	60	6	CQ545914	CQ545914 Sequence
C 23	58.2	6.6	9471	5	AY186728	AY186728 Gallus ga
C 24	58.2	6.6	125020	9	AF429315	AF429315 Homo sapi
C 25	57.4	6.5	82746	1	AF453501	AF453501 Actinosyn
C 26	57.2	6.4	3987	9	BC036533	BC036533 Homo sapi
C 27	57.2	6.4	300100	1	SC093123	AL939123 Streptomy
C 28	57	6.4	1822	5	GCCHX14	Y16952 Amycolatops
C 29	56.8	6.4	66669	1	AME16952	AL939106 Streptomy
C 30	56.8	6.4	314100	1	SC093106	AL939106 Streptomy
C 31	56.4	6.4	963	11	PM12H123	AL684456 Penicilli
C 32	55.6	6.3	943	9	AB042637	AB042637 Homo sapi
C 33	55.6	6.3	2507	9	AB042636	AB042636 Homo sapi
C 34	55.6	6.3	2811	9	AK055486	AK055486 Homo sapi
C 35	55.6	6.3	4023	9	AK126663	AK126663 Homo sapi
C 36	55.6	6.3	97110	2	AC016815	AC016815 Homo sapi
C 37	55.6	6.3	137960	2	AC103889	AC103889 Homo sapi
C 38	55.6	6.3	168133	2	AC093622	AC093622 Homo sapi
C 39	55.6	6.3	208236	6	AC010536	AC010536 Homo sapi
C 40	55.2	6.2	1265	8	AK108845	AK108845 Oryza sat
C 41	55.2	6.2	2997	6	AX552616	AX552616 Sequence
C 42	55.2	6.2	3158	6	AX552614	AX552614 Sequence
C 43	55.2	6.2	99575	2	AP004588	AP004588 Oryza sat
C 44	55.2	6.2	140142	8	AP005871	AP005871 Oryza sat
C 45	55.2	6.2	144879	8	AP005405	AP005405 Oryza sat

ALIGNMENTS

AF326964 887 bp mRNA linear PRI 01-MAY-2001
Homo sapiens BCLB (BCLB) mRNA, complete cds.
AF326964.1 GI:13898393
Homo sapiens (human)
ORGANISM
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
21201065
11278245
Ke.N., Godzik,A. and Reed,J.C.
Bcl-2, a novel Bcl-2 family member that differentially binds and
regulates Bax and Bak
J. Biol. Chem. 276 (16), 12481-12484 (2001)
REFERENCE
AUTHORS
TITLE
JOURNAL
MEDLINE
PUBMED
21201065
11278245
Ke.N., Godzik,A. and Reed,J.C.
Direct Submission
Submitted (07-DEC-2000) The Burnham Institute, 10901 N. Torrey
Pines Rd., La Jolla, CA 92037, USA
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/mol_type="mRNA"
/db_xref="taxon:9606"
/chromosome="15"
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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 5.4e-144;
Matches 887; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 CGGSCCAAGAAACACAGCGAAGCCCGGCCCCCAGAGAGCGGACCATGTTGACCA 60
Db 1 CGGSCCAAGAAACACAGCGAAGCCCGGCCCCCAGAGAGCGGACCATGTTGACCA 60

Qy 61 GTTGGGGAGCGGACACACATGCGCGGAGCCGCTGCGGGAGCGGACCGAGCTGTGCTGGC 120
Db 61 GTTGGGGAGCGGACACACATGCGCGGAGCCGCTGCGGGAGCGGACCGAGCTGTGCTGGC 120

Qy 121 CGACTACCTGGGTACTGCGCCCGGGAACCCGCGACCCCGGAGCGGCGCATCCACGCG 180
Db 121 CGACTACCTGGGTACTGCGCCCGGGAACCCGCGACCCCGGAGCGGCGCATCCACGCG 180

Qy 181 CGAGGCGCGCTGCTGCGCTCCGCGCCCGCCAGGTTAOGCAGANTCACGGTCTCTTTT 240
Db 181 CGAGGCGCGCTGCTGCGCTCCGCGCCCGCCAGGTTAOGCAGANTCACGGTCTCTTTT 240

Qy 241 CTCGGCTACTCGGCTACCCCGGAGACCGCTTCGAGCTGTGGCGCTGATGCGGATTC 300
Db 241 CTCGGCTACTCGGCTACCCCGGAGACCGCTTCGAGCTGTGGCGCTGATGCGGATTC 300

Qy 301 CGTGCTCTCGGACAGCCCGGCCCCCAGTGGGAGAGTGTGACGCTCGTGACCTTCGC 360
Db 301 CGTGCTCTCGGACAGCCCGGCCCCCAGTGGGAGAGTGTGACGCTCGTGACCTTCGC 360

Qy 361 AGGAGCCTGCTGAGAGAGCGCGCTGTGTGACCGCGCGTGAAGAAGTGGGCTTCCA 420
Db 361 AGGAGCCTGCTGAGAGAGCGCGCTGTGTGACCGCGCGTGAAGAAGTGGGCTTCCA 420

Qy 421 GCCCGGCTAAAGAGAGAGGAGCGGACGCTGCGCCCGGAGTGCACGCGCTGTGGCTTT 480
Db 421 GCCCGGCTAAAGAGAGAGGAGCGGACGCTGCGCCCGGAGTGCACGCGCTGTGGCTTT 480

Qy 481 GCTGAGCTCGGCGCTCATGCGGAGACCGCGCTGCTGAGGCTCAGGCGGCTGGGA 540
Db 481 GCTGAGCTCGGCGCTCATGCGGAGACCGCGCTGCTGAGGCTCAGGCGGCTGGGA 540

Qy 541 TGGCTTTTGTCACTTTTCAAGGACCCCTTTTCACTGGCTTTTGGAGAAACAGCTGCT 600
Db 541 TGGCTTTTGTCACTTTTCAAGGACCCCTTTTCACTGGCTTTTGGAGAAACAGCTGCT 600

Qy 601 CCAGGCTTTTCTGTGCTGTGTAAACACAGCGCTTCATTTATCTGTGACACGATATT 660
Db 601 CCAGGCTTTTCTGTGCTGTGTAAACACAGCGCTTCATTTATCTGTGACACGATATT 660

Qy 661 ATGAGTTTAAACTTTTAACTTTTAACTTTTAACTTTTAACTTTTAACTTTTAACTTT 720
Db 661 ATGAGTTTAAACTTTTAACTTTTAACTTTTAACTTTTAACTTTTAACTTTTAACTTT 720

Qy 721 ATGTGTGAGAACAGAACTGAGGAAAGACCTTCCCGACCCCGACGCTTTTATCTGA 780
Db 721 ATGTGTGAGAACAGAACTGAGGAAAGACCTTCCCGACCCCGACGCTTTTATCTGA 780

Qy 781 ATGCATACAGGAGTCTGTGAGTGTGATTTGGCGGAGTGTAACTTGTGACAGTACT 840
Db 781 ATGCATACAGGAGTCTGTGAGTGTGATTTGGCGGAGTGTAACTTGTGACAGTACT 840

Qy 841 CAGGTGTGAGGACAAAGATGCAATGCTCTTCTTGTAGTGAAGAA 887
Db 841 CAGGTGTGAGGACAAAGATGCAATGCTCTTCTTGTAGTGAAGAA 887

RESULT 2

BD233466

LOCUS

DEFINITION

ACCESSION

BD233466 1168 bp DNA linear PAT 17-JUL-2003
Human protein having hydrophobic domain and DNA encoding the same.
BD233466

VERSION

BD233466.1 GI:33043236

KEYWORDS

JP 2002519016-A/12.

SOURCE

Homo sapiens (human)

ORGANISM

Homo sapiens

REFERENCE

1 (bases 1 to 1168)

AUTHORS

Kato, S. and Kimura, T.

TITLE

Human protein having hydrophobic domain and DNA encoding the same

JOURNAL

Patent: JP 2002519016-A 12 02-JUL-2002;

COMMENT

SAGAMI CHEMICAL RESEARCH CENTER, PROTEGENE INC

OS Homo sapiens (human)

PN JP 2002519016-A/12

PD 02-JUL-2002

PF 18-JUN-1999 JP 2000557267

PI SEISHI KATO, TOMOKO KIMURA

PC

C12N15/09, C07K14/47, C12N1/15, C12N1/19, C12N5/10, C12N15/00, C12N5/ PC

CC Human protein having hydrophobic domain and DNA encoding the

CC same

FH Key

FT source

Location/Qualifiers

1..1168

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/mol_type="genomic DNA"

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ORIGIN

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Best Local Similarity 99.9%; Pred. No. 4.5e-131;
Matches 813; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 74 ACCACCATGGCCGACCCGCTGCGGAGCGCACCGAGCTGTGTGGCCGACTACCTGGGG 133
Db 1 ACCACCATGGCCGACCCGCTGCGGAGCGCACCGAGCTGTGTGGCCGACTACCTGGGG 60

Qy 134 TACTGCGCCCGGGAAACCGCGACCCCGAGCGCGCCATCCACGCCCGAGCGCGCGGTG 193
Db 61 TACTGCGCCCGGGAAACCGCGACCCCGAGCGCGCCATCCACGCCCGAGCGCGCGGTG 120

Qy 194 CTGCGCTCCGCGCGCGCGCAGGTTACGGCAGATTACCGGTCCTTTTCTCCGCTACCTC 253
Db 121 CTGCGCTCCGCGCGCGCGCAGGTTACGGCAGATTACCGGTCCTTTTCTCCGCTACCTC 180

Qy 254 GGCTACCCCGGGAAACCGCTTCGAGCTGTGGCGCTGATGCGGATTCGCTCTCCGAC 313
Db 181 GGCTACCCCGGGAAACCGCTTCGAGCTGTGGCGCTGATGCGGATTCGCTCTCCGAC 240

Qy 314 AGCCCGCGCCACCTCGGGCAGAGTGTGACGCTCGTGACCTTCGAGGGAAGCTGCTG 373
Db 241 AGCCCGCGCCACCTCGGGCAGAGTGTGACGCTCGTGACCTTCGAGGGAAGCTGCTG 300

Qy 374 CAGAGAGGCGCGCTGTGTGACCGCCCGGTGGAAGTGGGGCTTCAGCCCGCGCTAAAG 433
Db 301 CAGAGAGGCGCGCTGTGTGACCGCCCGGTGGAAGTGGGGCTTCAGCCCGCGCTAAAG 360

Qy 434 CAGCAGAGGCGCGACGCTGCGCCGGGACTGCCAGCGCTGTGGCCTTGCTGAGCTCCGG 493
Db 361 CAGCAGAGGCGCGACGCTGCGCCGGGACTGCCAGCGCTGTGGCCTTGCTGAGCTCCGG 420

Qy 494 CTCATGGGCGCAGCACCGCGCTGCTCAGGCTCAGGCGCGCTGGGATGCTTTTGTAC 553
Db 421 CTCATGGGCGCAGCACCGCGCTGCTCAGGCTCAGGCGCGCTGGGATGCTTTTGTAC 480

Qy 554 TTCCTCAGGACCCCTTTTCCACTGGCTTTTGGAGAAACAGCTGGTCCAGGCTTTTCTG 613
Db 481 TTCCTCAGGACCCCTTTTCCACTGGCTTTTGGAGAAACAGCTGGTCCAGGCTTTTCTG 540

Qy 614 TCATGCTGTTTAAACACAGCGCTTCATTATCTCTGGACAGATTATTATGAGTTTAAAA 673

Db	541	TCATGCTGTTTAAACACAGCCTTCATTTATCTCTGGACAGTATTATTAGTATTTAAAA	600
Qy	674	CTTTTAAACCGGCTTTCACCTGCGCCAACTGTGACCACTAAATGACAGATGTGTGAGAACA	733
Db	601	CTTTTAAACCGGCTTTCACCTGCGCCAACTGTGACCACTAAATGACAGATGTGTGAGAACA	660
Qy	734	AGAACTGAGGGAAGAACACCTTCCCAACCCAGAGAGTATTTATCTGAATGATCAAGAA	793
Db	661	AGAACTGAGGGAAGAACACCTTCCCAACCCAGAGAGTATTTATCTGAATGATCAAGAA	720
Qy	794	GTCTCTGAGTGTGGTGAATTTGGCCAGTGTATTTACTTGTGACAAAGTACTCAGGTGTGAGGAC	853
Db	721	GTCTCTGAGTGTGGTGAATTTGGCCAGTGTATTTACTTGTGACAAAGTACTCAGGTGTGAGGAC	780
Qy	854	AGAAATGCAAAATGGCTCTTCCTTGAAGTGAAGAA	887
Db	781	AGAAATGCAAAATGGCTCTTCCTTGAAGTGAAGAA	814
RESULT 3			
AF285092	AF285092	615 bp mRNA	linear PRI 08-NOV-2001
LOCUS	Homo sapiens Bcl-2-like protein 10 mRNA, complete cds.		
DEFINITION	AF285092		
ACCESSION	AF285092.1	GI:9837265	
VERSION			
KEYWORDS	Homo sapiens (human)		
SOURCE	Homo sapiens		
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
REFERENCE	1 (bases 1 to 615)		
AUTHORS	Zhang, H., Holzgrevé, W. and De Geyter, C.		
TITLE	Bcl2-L-10, a novel anti-apoptotic member of the Bcl-2 family, blocks apoptosis in the mitochondria death pathway but not in the death receptor pathway		
JOURNAL	Hum. Mol. Genet. 10 (21), 2329-2339 (2001)		
MEDLINE	21548034		
PUBMED	11689480		
REFERENCE	2 (bases 1 to 615)		
AUTHORS	Zhang, H. H.		
TITLE	Direct Submission		
JOURNAL	Submitted (05-JUL-2000) University Women's Hospital, Schanzenstr		
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Best Local Similarity 100.0%; Pred. No. Se-97;			
Matches 615; Conservative 0; Mismatches 0; Indels 0; Gaps 0;			
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Db	1	ATGGTTGACAGTTGGGGAGCGCACACCATGGCCGACCCCTCGGGAGCGCACCCGAG	60
Qy	110	CTGTTCTGCGCGACTACTCTGGGGTACTCGCCCGGGAAACCCGGACCCCGAGCCGCG	169
Db	61	CTGTTCTGCGCGACTACTCTGGGGTACTCGCCCGGGAAACCCGGACCCCGAGCCGCG	120

Qy	170	CCATCCACGCCGAGGCGCGCTGCTCGCTCCGCGCCGCGCGCTTACGCGAGATTAC	229
Db	121	CCATCCACGCCGAGGCGCGCTGCTCGCTCCGCGCCGCGCGCTTACGCGAGATTAC	180
Qy	230	CGGTCCTTTTCTCCGCTACCTCGGCTACCCCGGAGAACCGCTTCGAGCTGGTGGGCTG	289
Db	181	CGGTCCTTTTCTCCGCTACCTCGGCTACCCCGGAGAACCGCTTCGAGCTGGTGGGCTG	240
Qy	290	ATGCGGATTCGCGCTCTCCGACAGCCCGCGCCCACTTGGGGCAGAGTGGTGACGCTC	349
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Qy	350	GTGACCTTTCGAGGAGCGCTGCTGGAGAGGCGCGCTGTGACCCGCCCGTGAAGAAG	409
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Qy	410	TGGGCTTCCAGCGCGCTAAAGGAGCAGAGGAGCGCTGCCCGGGACTGCCAGCGC	469
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Qy	470	CTGTGCGCTTGTGAGCTCGCGCTCATGGGAGCAGCACCGCGCTGGTGCAGGCTCAG	529
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Qy	530	GGCGGCTGGGATGGCTTTTGTCTCACTTTTCAGGACCCCTTCCACTGGGCTTTTGAGA	589
Db	481	GGCGGCTGGGATGGCTTTTGTCTCACTTTTCAGGACCCCTTCCACTGGGCTTTTGAGA	540
Qy	590	AAACAGCTGTGCCAGGCTTTTCTGTGATGTTGTTAAACACAGCCTTCAATTTCTCTGG	649
Db	541	AAACAGCTGTGCCAGGCTTTTCTGTGATGTTGTTAAACACAGCCTTCAATTTCTCTGG	600
Qy	650	ACACGATTATATGA	664
Db	601	ACACGATTATATGA	615
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DEFINITION	Homo sapiens NRH gene for anti-apoptotic protein.		
ACCESSION	AJ458330		
VERSION	AJ458330.1	GI:20338765	
KEYWORDS	anti-apoptotic protein; NRH gene.		
SOURCE	Homo sapiens (human)		
ORGANISM	Homo sapiens		
REFERENCE	1	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.	
AUTHORS	1	Aouacheria, A., Arnaud, E., Venet, S., Lalle, P., Gouy, M., Rigal, D., and Gillet, G.	
TITLE	Nrh, a human homologue of Nr-13 associates with Bcl-Xs and is an inhibitor of apoptosis		
JOURNAL	Oncogene 20 (41), 5846-5855 (2001)		
MEDLINE	21477277		
PUBMED	11593390		
REFERENCE	2 (bases 1 to 585)		
AUTHORS	Gillet, G.		
TITLE	Direct Submission		
JOURNAL	Submitted (23-APR-2002) Gillet G., Ibcop, CNRS UMR 5086, 7 PASSAGE DU VERCORS, 69367, FRANCE		
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CDS			

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ORIGIN

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Best Local Similarity 100.0%; Pred. No. 1.3e-91;
Matches 585; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 80 ATGGCCGACCCGCTGGGGAGCCACCGAGCTGTGCTGGCCGACTACCTGGGGTACTGC 139
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QY 140 GCCCGGGAACCCGGACCCCGAGCCGCGCCATCCACGCCCGAGGCCCGCTGTGGCC 199
DB 61 GCCCGGGAACCCGGACCCCGAGCCGCGCCATCCACGCCCGAGGCCCGCTGTGGCC 120

QY 200 TCCGCGCGCCAGGTTACCGGAGATTACCGGTCTCTTTTCTCGCCCTACCTCGGCTAC 259
DB 121 TCCGCGCGCCAGGTTACCGGAGATTACCGGTCTCTTTTCTCGCCCTACCTCGGCTAC 180

QY 260 CCCGGGAACCCGCTTCGAGCTGTGGCGCTGATGGCGGATTCGCTGTCTCGACAGCCCC 319
DB 181 CCCGGGAACCCGCTTCGAGCTGTGGCGCTGATGGCGGATTCGCTGTCTCGACAGCCCC 240

QY 320 GGGCCGCGCTGGGGAGAGTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 379
DB 241 GGGCCGCGCTGGGGAGAGTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 300

QY 380 GGGCCGCGCTGGGGAGAGTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 439
DB 301 GGGCCGCGCTGGGGAGAGTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 360

QY 440 GAGGGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 499
DB 361 GAGGGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 420

QY 500 GGGCAGCACCAGCCCTGGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 559
DB 421 GGGCAGCACCAGCCCTGGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 480

QY 560 AGGACCCCTTTTCCACTGGCTTTTGGAGAAACAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 619
DB 481 AGGACCCCTTTTCCACTGGCTTTTGGAGAAACAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 540

QY 620 TTGTTAAACAACAGCTTCATTATCTCTGACACGATTATTATGA 664
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RESULT 5

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BD233456
LOCUS 582 bp DNA linear PAT 17-JUL-2003
DEFINITION Human protein having hydrophobic domain and DNA encoding the same.
ACCESSION BD233456
VERSION BD233456.1 GI:33043226
KEYWORDS JP 2002519016-A/2.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 582)
Kato, S. and Kimura, T.
Human protein having hydrophobic domain and DNA encoding the same
Patent: JP 2002519016-A 2 02-JUL-2002;
SAGAMI CHEMICAL RESEARCH CENTER, PROTEGENE INC

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COMMENT OS Homo sapiens (human)
PN JP 2002519016-A/2
PD 02-JUL-2002
PF 18-JUN-1999 PF 2000557267
PI SEISHI KATO, TOMOKO KIMURA
PC C12N15/09, C07K14/47, C12N1/15, C12N1/19, C12N5/10, C12N15/00, C12N5/ PC
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CC Human protein having hydrophobic domain and DNA encoding the
CC same
FT Key Location/Qualifiers
FT source 1..582
FT /organism="Homo sapiens"
FT /mol_type="genomic DNA"
FT /db_xref="taxon:9606"

FEATURES
source
ORIGIN
Query Match 65.6%; Score 582; DB 6; Length 582;
Best Local Similarity 100.0%; Pred. No. 4.2e-91;
Matches 582; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 80 ATGGCCGACCCGCTGGGGAGCCACCGAGCTGTGCTGGCCGACTACCTGGGGTACTGC 139
DB 1 ATGGCCGACCCGCTGGGGAGCCACCGAGCTGTGCTGGCCGACTACCTGGGGTACTGC 60

QY 140 GCCCGGGAACCCGGACCCCGAGCCGCGCCATCCACGCCCGAGGCCCGCTGTGGCC 199
DB 61 GCCCGGGAACCCGGACCCCGAGCCGCGCCATCCACGCCCGAGGCCCGCTGTGGCC 120

QY 200 TCCGCGCGCCAGGTTACCGGAGATTACCGGTCTCTTTTCTCGCCCTACCTCGGCTAC 259
DB 121 TCCGCGCGCCAGGTTACCGGAGATTACCGGTCTCTTTTCTCGCCCTACCTCGGCTAC 180

QY 260 CCCGGGAACCCGCTTCGAGCTGTGGCGCTGATGGCGGATTCGCTGTCTCGACAGCCCC 319
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DB 301 GGGCCGCGCTGGGGAGAGTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAGCTGTGAG 360

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RESULT 6

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LOCUS 726 bp DNA linear PAT 03-FEB-2004
DEFINITION Sequence 38039 from Patent WO02068579.
ACCESSION CQ752105
VERSION CQ752105.1 GI:42387450
KEYWORDS Homo sapiens (human)
SOURCE

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ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1
REFERENCE Venter C.J., Adams M.C., Li P.W. and Myers E.W.
AUTHORS Kits, such as nucleic acid arrays, comprising a majority of
TITLE humanexons or transcripts, for detecting expression and other uses
thereof
JOURNAL Patent: WO 02068579-A 38039 06-SEP-2002;
PE Corporation (NY) (US)
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QY 170 CCATCCACCCCGAGCGCCGCTGTGCTCGCGCGCGCCAGCTTACGGCAGATTCCAC 229
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QY 230 CGGTCTCTTTCTCCGCTACTCTCGCTACCCCGGGAACCGTTGAGCTGTGGCGT 289
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RESULT 7
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LOCUS
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sequence.
ACCESSION AC023906
VERSION AC023906.7 GI:14595770
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 93287)
AUTHORS Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B.,
Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G.,
James,R., Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T.
and Hood,L.
Pate,D. and Hood,L.
TITLE Sequencing of human chromosome 15 D15S146-D15S117 region
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 93287)
AUTHORS Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B.,
Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G.,
James,R., Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T.
and Hood,L.
Pate,D. and Hood,L.
TITLE Direct Submission
JOURNAL Submitted (20-FEB-2000) Multimegabase Sequencing Center, University
of Washington, PO BOX 357730, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 93287)
AUTHORS Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B., Bloom,S.,
Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R.,
Pate,D. and Hood,L.
TITLE Direct Submission
JOURNAL Submitted (04-JUL-2001) Multimegabase Sequencing Center, Institute
for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA
98105, USA
REFERENCE 4 (bases 1 to 93287)
AUTHORS Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B., Bloom,S.,
Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R.,
Pate,D. and Hood,L.
TITLE Direct Submission
JOURNAL Submitted (06-JUL-2001) Multimegabase Sequencing Center, Institute
for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA
98105, USA
COMMENT On Jul 4, 2001 this sequence version replaced gi:12248292.

----- Genome Center
Center: Multimegabase Sequencing Center
Center code: UWMSC
Web site: http://chroma.mbt.washington.edu/msg_www
Contact: leerowens@systemsbiology.org
----- Summary Statistics
Sequencing vector: pUC18; L08752
Chemistry: Dye-terminator Big Dye; 90% of reads
Chemistry: Dye-primer Big Dye; 10% of reads
Assembly program: Phrap; version 0.990399

Note: Data from overlapping BACs AC010674 [drafting center:
UWMSC], AC090970 [drafting center: UWMSC]; and AC016824 [drafting
center: GTC] was added for finishing

FEATURES
Source

Location/Qualifiers

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/note="Data from overlapping clones CTD-2650P22 AC090970,
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consensus was determined from CTD-2184D3 to the extent
possible"

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Best Local Similarity 99.4%; Pred. No. 7.5e-84;
Matches 541; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
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Db 64931 GTTGGGGAGCGCACACCATGCGCCAGCCCGCTGCGGGAGCCACCGAGCTGTGCTGGC 64872
Qy 121 CGACTACTCGGGTACTGCGCCCGGAAACCGGCACCCCGGAGCCGCCATCCACGCC 180
Db 64871 CGACTACTCGGGTACTGCGCCCGGAAACCGGCACCCCGGAGCCGCCATCCACGCC 64812
Qy 181 CGAGGCGCGCTGCTGGGCTCGCGCCCGCAGGTTACGGCAGATTCACCGCTCTTTT 240
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SEQUENCE SAMPLING.
ACCESSION AC018903
VERSION AC018903.2 GI:8247797
KEYWORDS HTG; HTGS_PHASE0.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
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REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
1 (bases 1 to 214669)
Rowen, L., Madan, A., Qin, S., Abbasi, N., Baradarani, L., Birditt, B.,
Bloom, S., Dors, M., Dickhoff, R., Fleetwood, P., Harrison, G., Kaur, A.,
Madan, A., Nesbitt, R., Shaffer, T. and Hood, L.
Sequencing of human chromosome 15 D15S146-D15S117 region
Unpublished
2 (bases 1 to 214669)
Rowen, L., Madan, A., Qin, S., Abbasi, N., Baradarani, L., Birditt, B.,
Bloom, S., Dors, M., Dickhoff, R., Fleetwood, P., Harrison, G.,
James, R., Kaur, A., Madan, A., Owen, M. P., Ratcliffe, A., Shaffer, T.
and Hood, L.
Direct Submission
Submitted (22-DEC-1999) Multimegabase Sequencing Center, University
of Washington, PO BOX 357730, Seattle, WA 98195, USA
On Jun 4, 2000 this sequence version replaced gi:6630517.
----- Genome Center
Center: Multimegabase Sequencing Center
Center code: UMWSC
Web site: http://chroma.mbt.washington.edu/msg_www
Contact: leerowens@systemsbiology.org
----- Summary Statistics
Sequencing vector: pUC18; 108752
Chemistry: Dye-terminator Big Dye; 90% of reads
Chemistry: Dye-primer Big Dye; 10% of reads
Assembly program: Phrap; version 0.990399

* NOTE: This record contains 192 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.

1
806: contig of 806 bp in length
906: gap of unknown length
907: contig of 1162 bp in length
2068: gap of unknown length
2069: contig of 852 bp in length
3020: gap of unknown length
3120: contig of 1164 bp in length
4284: gap of unknown length
4384: gap of unknown length
5201: contig of 817 bp in length
5301: gap of unknown length
5544: contig of 1243 bp in length
6644: gap of unknown length
7472: contig of 828 bp in length
7572: gap of unknown length
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13393: gap of unknown length
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15468: contig of 1154 bp in length
15565: gap of unknown length
16379: contig of 814 bp in length
16479: gap of unknown length
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Qy 656 TTATTATGAGCTTTTAAACCTTTTAAACCCGCTTCTACTGCCCACTGTGACCACTAAAT 715
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Qy 716 GACAGATGTGTGAGAACAGAACTGAGGGAAGAACACCTTCCCCACCCAGAGCTTTT 775
Db 168402 GACAGATGTGTGAGAACAGAACTGAGGGAAGAACACCTTCCCCACCCAGAGCTTTT 168343

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RESULT 9
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SEQUENCE SAMPLING.
ACCESSION
VERSION AC018903.2 GI:8247797
KEYWORDS HTG; HTGS PHASE0.
SOURCE
ORGANISM Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B.,
Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G., Kaur,A.,
Madan,A., Nesbitt,R., Shaffer,T. and Hood,L.
Sequencing of human chromosome 15 D15S146-D15S117 region
Unpublished
2 (bases 1 to 214669)
Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B.,
Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G.,
James,R., Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T.
and Hood,L.
Direct Submission
Submitted (22-DEC-1999) Multimegabase Sequencing Center, University
of Washington, PO BOX 357730, Seattle, WA 98195, USA
On Jun 4, 2000 this sequence version replaced gi:6630517.
----- Genome Center
Center: Multimegabase Sequencing Center
Web site: http://chroma.mbt.washington.edu/msg_www
Contact: leerowen@systemsbiology.org
----- Summary Statistics
Sequencing vector: pUC18; 108752
Chemistry: Dye-terminator Big Dye; 90% of reads
Chemistry: Dye-primer Big Dye; 10% of reads
Assembly program: Phrap; version 0.990399
-----
* NOTE: This record contains 192 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
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14312 15465: contig of 1154 bp in length
15466 16379: contig of 814 bp in length
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16480 17624: contig of 1145 bp in length
17625 17725: gap of unknown length
17726 18578: contig of 854 bp in length
18579 18679: gap of unknown length
18680 19774: contig of 1096 bp in length
19775 20700: contig of 826 bp in length
20701 20800: gap of unknown length
20801 22136: contig of 1236 bp in length
22137 23023: contig of 887 bp in length
23024 23123: gap of unknown length
23124 24196: contig of 1073 bp in length
24197 24296: gap of unknown length
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26386 26485: gap of unknown length
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27326 27425: gap of unknown length
27426 28691: contig of 1266 bp in length
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31914 33056: contig of 1143 bp in length
33057 34007: contig of 851 bp in length
34008 34107: gap of unknown length
34108 35320: contig of 1213 bp in length
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37522 37621: gap of unknown length
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39727 39826: gap of unknown length
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44926 46114: contig of 1189 bp in length
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48293 48392: gap of unknown length
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* 57659 57659: contig of 851 bp in length
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Matches 406; Conservative 0; Mismatches 94; Indels 2; Gaps 2;

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QY 61 GTTGGCGGAGCGCACCAACCATGCGGACCCGCTGCGGGAGCGCACCGAGCTGTTCTCTGGC 120
DB 93273 TTTGCGGAGCGCACCAACCATGTCGACCCGCTGCGGGAGCGCACCGAGCTGTTCTCTGGC 93332
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QY 300 CCGTGTCTCCGACAGCCCGCGCGCGCGCGAGTGGTGGCGAGTGGTGGCGAGTGGTGGCGCTTCG 359
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QY 360 CAGGAGCGCTGCTGGAGAGAGCGCCGCTGCTGACCCCGCGTGGAGAAAGTGGGGCTTC 419
DB 93573 CAGGACACTTCTTGAGAAAGGCCCTAGTAGCCCGCCCTTGAAAAAATAGCTTTC 93632
QY 420 AGCGCGGCTAAAGGAGCAGGAGGCGAGCTGCGCCCGGAGCTGCCAGCGCTGGTGGCTTC 479
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QY 480 TGCTGAGCTCGCGCTCATGGG 501
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DEFINITION AF102501
ACCESSION AF102501
VERSION AF102501.1 GI:4165137
KEYWORDS
SOURCE
ORGANISM
Mus musculus (house mouse)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 1209)
Song,Q., Kuang,Y., Dixit,V.M. and Vincenz,C.
Boo, a novel negative regulator of cell death, interacts with
Apaf-1
EMBO J. 18 (1), 167-178 (1999)
JOURNAL
MEDLINE 99094902
PUBMED 9878060
REFERENCE
2 (bases 1 to 1209)
Song,Q.Z., Kuang,Y.P., Dixit,V.M. and Vincenz,C.
Direct Submission
Submitted (28-OCT-1998) Pathology, University of Michigan, 1301
Catherine Road, Ann Arbor, MI 48109, USA
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Qy 206 GCGCCGAGGTTACGCGAGATTACCGGTCTTTTCTCGGCTACCTCGGCTACCCCGG 265
Db 252 ACTAGGAGATCAGCAGAGACCAAGAAATTTTCTCTCTCTGCAAGACCGGGGC 311
Qy 266 AACCGCTTCGAGCTGTGGCGCTGTGATGGCGATTCGCTCTCCGACAGCCCGCGCC 325
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Qy 326 ACTGGGCGAGCTGTGAGCTGTGACCTTCGACGAGGACGCTGTGGAGAGGGCG 385
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Qy 386 CTGGTGACCCCGCGGTGGAGAAAGTGGGGCTTCACGCCGCGCTAAAGGAGCAGAGGCG 445
Db 432 TACATGGCTGTCAAGCAGAAGA-----GGATCTGGGGAATCGTGTG 473
Qy 446 GAGTGCCTCGGAGCTGCGAGCCCTGTGGCTTCTGAGCTCGCGGCTCATGGG--- 502
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ACCESSION AF067660
VERSION AF067660.1 GI:3955265
KEYWORDS
SOURCE Mus musculus (house mouse)
ORGANISM
REFERENCE 1 (bases 1 to 1225)
AUTHORS Inohara, N., Gourley, T. S., Carrio, R., Muniz, M., Merino, J., Garcia, I., Koski, T., Hu, Y., Chen, S., and Nunez, G.
TITLE Diva, a Bcl-2 homologue that binds directly to Apaf-1 and induces BH3-independent cell death

J. Biol. Chem. 273 (49), 32479-32486 (1998)
99047617
PUBMED 9829980
REFERENCE 2 (bases 1 to 1225)
AUTHORS Inohara, N. and Nunez, G.
TITLE Direct Submission
JOURNAL Submitted (20-MAY-1998) Department of Pathology, University of Michigan Medical School, 1500 E. Medical Dr., Ann Arbor, MI 48109, USA
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Matches 477; Conservative 0; Mismatches 253; Indels 26; Gaps 4;
Qy 86 GACCCGCTGGGAGCGCACCGAGCTGTTGCTGGCGACTACTCTGGGTACTGGCGCCGG 145
Db 158 GACCCACTGATGAACGCACTAGACGGCTGCTGTCTGACTACATATTCTTCTGCGACGG 217
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Db 398 AGCTGGAGCCCACTGGTGTCTCTTCTGCGGAGCGCTTATGAATCAAGCCCT 457
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LOCUS
DEFINITION
Mus musculus Bcl2-like 10, mRNA (cDNA clone MGC:60542
IMAGE:30052580), complete cds.
ACCESSION
BC052690
VERSION
MGC.
KEYWORDS
SOURCE
Mus musculus (house mouse)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 1257)
Strausberg,R.L., Feingold,E.A., Grouse,L.H., Derge,J.G.,
Klausner,R.D., Collins,F.S., Wagner,L., Shenmen,C.M., Schuler,G.D.,
Altschul,S.F., Zeeberg,B., Buetow,K.H., Schaefer,C.F., Bhat,N.K.,
Hopkins,R.F., Jordan,H., Moore,T., Max,S.I., Wang,J., Hsieh,F.,
Diatchenko,L., Marusina,K., Farmer,A.A., Rubin,G.M., Hong,L.,
Staatelen,M., Soares,M.B., Bonaldo,M.F., Casavant,T.L.,
Schetz,T.E., Brownstein,M.J., Usdin,T.B., Toshiyuki,S.,
Carninci,P., Frange,C., Raha,S.S., Loquellano,N.A., Peters,G.J.,
Abramson,R.D., Mullaly,S.J., Bosak,S.A., McEwan,P.J.,
McKernan,K.J., Malek,J.A., Gunaratne,P.H., Richards,S.,
Worley,K.C., Hale,S., Garcia,A.M., Gay,L.J., Rulyk,S.W.,
Villalon,D.K., Muzny,D.M., Sodergren,E.J., Lu,X., Gibbs,R.A.,
Fahney,J., Hellon,E., Kettman,M., Madan,A., Rodrigues,S.,
Sanchez,A., Whitting,M., Madan,A., Young,A.C., Shevchenko,Y.,
Bouffard,G.G., Blakesley,R.W., Touchman,J.W., Green,E.D.,
Dickson,M.C., Rodriguez,A.C., Grimwood,J., Schmutz,J., Myers,R.M.,
Butterfield,Y.S., Krzywinski,M.I., Skalska,U., Smalls,D.E.,
Schurch,A., Schein,J.E., Jones,S.J. and Marra,M.A.
Generation and initial analysis of more than 15,000 full-length
human and mouse cDNA sequences
Proc. Natl. Acad. Sci. U.S.A. 99 (26), 16899-16903 (2002)
22388257
12477932
2 (bases 1 to 1257)
Strausberg,R.
Direct Submission
Submitted (16-MAY-2003) National Institutes of Health, Mammalian
Gene Collection (MGC), Cancer Genomics Office, National Cancer
Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590,
USA
NIH-MGC Project URL: http://mgc.nci.nih.gov
Contact: MGC help desk
Email: cgabs-r@mail.nih.gov
Tissue Procurement: Minoru Ko
cDNA Library Preparation: Yulan Piao and Minoru Ko (National
Institute on Aging, NIH: http://lgsun.grc.nia.nih.gov/cDNA/)
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LINL)
DNA Sequencing by: National Institutes of Health Intramural
Sequencing Center (NISC),
Gaithersburg, Maryland;
Web site: http://www.nisc.nih.gov/
Contact: nisc_mgc@nhgri.nih.gov

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Akhter, N., Ayele, K., Beckstrom-Sternberg, S.M., Benjamin, B., Blakesley, R.W., Bouffard, G.G., Breen, K., Brinkley, C., Brooks, S., Dietrich, N.L., Granite, S., Guan, X., Gupta, J., Haghighi, P., Hansen, N., Ho, S.-L., Karlins, E., Kwong, P., Laric, P., Legaspi, R., Maduro, Q.L., Maselli, C., Maskeri, B., Mastrian, S.D., McCloskey, J.C., McDowell, J., Pearson, R., Stantiripop, S., Thomas, P.J., Touchman, J.W., Tsurgeon, C., Vogt, J.L., Walker, M.A., Wetherby, K.D., Wiggins, L., Young, A., Zhang, L.-H. and Green, E.D.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LINL at: <http://image.llnl.gov>
 Series: IRAC Plate: 112 Row: h Column: 18
 This clone was selected for full length sequencing because it passed the following selection criteria: matched mRNA gi: 7304926.

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QY 86 GACCCGCTGGGAGCGCACCAGCTGTGCTGGCGACTTACCTGGGTAAGTCTGGCCGG 145
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 LOCUS Rattus norvegicus BCL2L10 (Bcl2l10) mRNA, complete cds.

ACCESSION AY029163
 VERSION AY029163.1 GI:13641257
 KEYWORDS
 SOURCE Rattus norvegicus (Norway rat)
 ORGANISM Rattus norvegicus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae;
 Rattus.

REFERENCE 1 (bases 1 to 1074)
 Itoh,T., Itoh,A. and Pleasure,D.
 Bcl-2-related protein family gene expression during
 oligodendroglial differentiation
 J. Neurochem. 85 (6), 1500-1512 (2003)
 MEDLINE 22672518
 PUBMED 12787069
 REFERENCE 2 (bases 1 to 1074)
 Itoh,T., Itoh,A. and Pleasure,D.
 Direct Submission
 Submitted (29-NAR-2001) Neurology Research, The Children's Hospital
 of Philadelphia, Abramson Research Center, Room 516 I, 3517 Civic
 Center Boulevard, Philadelphia, PA 19104, USA
 Location/Qualifiers

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CDS

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ORIGIN

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 Best Local Similarity 61.7%; Pred. No. 1.5e-32;
 Matches 455; Conservative 0; Mismatches 253; Indels 29; Gaps 3;
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 Db 12 ATGGGTGACCCGCTGAGGATCCACTAGCGGCTGCTGACTACTACATATGTTCTGC 71
 QY 140 GCCCGGAACCCCGCACCCCGGAGCGGCCCATCCAGCCCGGCGCGCTGCTGCGC 199
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 QY 200 TCCGCGCCCGCAGGTTCAGGAGATTACCGGCTCTCTTTTCTCCGCTACCTCGGCTAC 259
 Db 132 TCTGTGACTAGTCAGATCCCAACAGGAGCACAGATCTTTTCAACTCTTCCGCGACTAC 191
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 Db 192 CAGGCGAACCCCTGGAGCTGGTGACACAGATGGCGGATGAGTTGCTTCCATGACCAA 251
 QY 320 GGGCCCACTGGGCGCAGATGGTGAAGCTGTCACCTTCCAGGAGCGTCTGGAGAGA 379
 Db 252 GAGTTCAACTGGGCGCGCTGGTGTGCTCTCGGCTTCTGGGAGCGTAAATGAACCAA 311
 QY 380 GGGCGCTGTGTGACCGCGCCGCTGGAGAGTGGGGCTTCCAGCGCGGCTAAAGGAGCAG 439
 Db 312 GACAGGACTGTTAAGCGGAGGAGGATCAAGAAACCGTCTCTCTAC----- 357
 QY 440 GAGGCGACGCTCGCCCGGAGCTGCGAGCGCTGGTGGCTTGTGAGCTCGCGGCTCATG 499
 Db 358 -----TGGAGCGAGACTGCTATCTCATAGTGAAGTGTGTACAAATCGACTCACA 407
 QY 500 GGGCAGCAGCGCGCTGGCTGCGAGGCTCAGGCGGCTGGGATGGCTTTTGTCACTTCTTC 559
 Db 408 GGACGGCATGCTCTCTGGCTGGAGGCTCAGGCTGGGATGGCTTTTGGCAATCTTC 467
 QY 560 AGGACCCCTTTCACATGGCTTTTGGAGAAACAGCTGTCCAGGCTTTTCTGTATGC 619
 Db 468 AAGAACCCCTTACACCCCGCTTCTGGAGAAAGTGTGATCCGGGCTATTTCTGCTGT 527
 QY 620 TTGTTAAACAGCGCTTCATTTATCTCTGGACAGATTATATGAGTTTAAACTTTTA 679
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 QY 680 ACCCGCTTCTACCTGCCCAACTGTGACCAACTAAATGACAGATGTGTGAGAACAGAACT 739
 Db 584 AATCGATCTTACCTGCTCTACTGCGGCCCTCTTAAGGGACAATTTGGTGTAGTGATAA 643
 QY 740 GAGGGAAGACCTTCCCGCAGCGCTTTTATCTGAATGCATACAGGAGCTCTG 799
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RESULT 14

AC133947

LOCUS

DEFINITION

Mus musculus chromosome 9 clone RP24-18912, WORKING DRAFT SEQUENCE,
 3 unordered pieces.

ACCESSION

AC133947

VERSION

AC133947.2 GI:50199129

KEYWORDS

HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ACTIVEFIN.

SOURCE

Mus musculus (house mouse)

ORGANISM

Mus musculus

AC133947 169914 bp DNA linear HTG 10-JUL-2004
 Mus musculus chromosome 9 clone RP24-18912, WORKING DRAFT SEQUENCE,
 3 unordered pieces.

ACCESSION AC133947
 VERSION AC133947.2 GI:50199129
 KEYWORDS HTG; HTGS PHASE1; HTGS DRAFT; HTGS_ACTIVEFIN.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 169914)
Wilson, R.K.
The sequence of Mus musculus clone
Unpublished
2 (bases 1 to 169914)
McPherson, J.D. and Waterston, R.H.
Direct Submission
Submitted (20-SEP-2002) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
3 (bases 1 to 169914)
Wilson, R.K.
Direct Submission
Submitted (10-JUL-2004) Genome Sequencing Center, 4444 Forest Park
Parkway, St. Louis, MO 63108, USA
On Jul 10, 2004 this sequence version replaced gi:23238076.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu
Contact: submissions@wustl.wustl.edu
----- Project information -----
Center project name: M_BB0189102
----- Summary Statistics -----
Sequencing vector: M13; 0%
Sequencing vector: plasmid; 100%
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 167792 bases at least Q40
Consensus quality: 168148 bases at least Q30
Consensus quality: 168291 bases at least Q20

* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence.
* as soon as it is available and the accession number will
* be preserved.

1 21204: contig of 21204 bp in length
* 21205 21304: gap of unknown length
* 21305 86045: contig of 64741 bp in length
* 86046 86145: gap of unknown length
* 86146 169914: contig of 83769 bp in length.

FEATURES

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ORIGIN

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Best Local Similarity 61.5%; Pred. No. 7e-16;
Matches 281; Conservative 0; Mismatches 155; Indels 21; Gaps 2;
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DB 83475 GACCCACTGCATGAAGCGACTAGACGGTGTGTGACTACATATTTCTTCGCGCAGG 83534

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LOCUS AC115880 202851 bp DNA linear ROD 02-APR-2004
DEFINITION Mus musculus chromosome 9, clone RP24-365N15, complete sequence.
ACCESSION AC115880
VERSION AC115880.11 GI:46063808
KEYWORDS HTG.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE
1 (bases 1 to 202851)
Barren, B., Nusbaum, C. and Lander, E.
Mus musculus chromosome 9, clone RP24-365N15
Unpublished
2 (bases 1 to 202851)
Barren, B., Linton, L., Nusbaum, C., Lander, E., Ali, A., Allen, N.,
Anderson, S., Barna, N., Bastien, V., Bloom, T., Boguslavsky, L.,
Boukhalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J.,
Chazaro, B., Choepel, Y., Colangelo, M., Collins, S., Collymore, A.,
Cook, A., Cooke, P., DeArellano, K., Dewar, K., Diaz, J.S., Dodge, S.,
Faro, S., Ferreira, P., FitzHugh, W., Gage, D., Galagan, J., Gardyna, S.,
Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
Kamat, A., Karatas, A., Kells, C., Larocque, K., Lamazares, R.,
Landers, R., Lehoczy, J., Levine, R., Lindblad-Toh, K., Liu, G.,
MacLean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C.,
McCarthy, M., McSwan, P., McKernan, K., Meldrim, J., Meneus, L.,
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Norbu, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D.,
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Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Scojanovic, N.,
Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G.,
Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE

Direct Submission
Submitted (22-MAR-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE
3 (bases 1 to 202851)
Barren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N.,

Anderson, M., Arachchi, H.M., Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L., Boukhalter, B., Camarata, J., Chang, J., Choepel, Y., Collamore, A., Cook, A., Cooke, P., Corum, B., DeAtellano, K., Diaz, J.S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, X., Lui, A., Mabbitt, R., MacLean, C., Macdonald, P., Major, J., Manning, J., Matthews, C., McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Rachupka, A., Ramaeamy, U., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schuback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R., Vo, A., Wilson, B., Wu, X., Wymann, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission
Submitted (28-FEB-2004) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 202851)

TITLE JOURNAL

REFERENCE AUTHORS

Anderson, M., Arachchi, H.M., Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L., Boukhalter, B., Camarata, J., Chang, J., Choepel, Y., Collamore, A., Cook, A., Cooke, P., Corum, B., DeAtellano, K., Diaz, J.S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J., Gardyna, S., Graham, L., Grand-Pierre, N., Hafez, N., Hagopian, D., Hagos, B., Hall, J., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, X., Lui, A., Mabbitt, R., MacLean, C., Macdonald, P., Major, J., Manning, J., Matthews, C., McCarthy, M., Meldrim, J., Meneus, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nguyen, T., Nicol, R., Norbu, C., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Rachupka, A., Ramasamy, U., Raymond, C., Retta, R., Rise, C., Rogov, P., Roman, J., Schauer, S., Schuback, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Stubbs, M., Talamas, J., Tesfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H., Venkataraman, V.S., Viel, R., Vo, A., Wilson, B., Wu, X., Wymann, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

Direct Submission
Submitted (02-APR-2004) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Apr 2, 2004 this sequence version replaced gi:44681569.
All repeats were identified using RepeatMasker:
Smit, A.P.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RN/RepeatMasker.html>

TITLE JOURNAL

COMMENT

Center: Whitehead Institute/MIT Center for Genome Research
Web site: <http://www.seq.wi.mit.edu>
Contact: sequence_submissions@broad.mit.edu
----- Project Information
Center project name: L24773
Center clone name: 365_N15

FEATURES source

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Query Match 16.6%; Score 147; DB 10; Length 202851;
Best Local Similarity 61.5%; Pred.No. 6.9e-16;
Matches 281; Conservative 0; Mismatches 155; Indels 2; Gaps 2;

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Job time : 6240 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: November 7, 2004, 03:03:17 ; Search time 510 Seconds

Title: US-10-071-174-2
Perfect score: 1084
Sequence: 1 MVDQLERTTMADPLRTERE.....QAFSLCLTTAFIVLWTELL 204

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Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 4134886 seqs, 2624710521 residues

Total number of hits satisfying chosen parameters: 8269772

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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5	639	58.9	548	ABK41913	Abk41913 cDNA enco
6	639	58.9	548	ADB59580	Adb59580 Connectiv

7	559	51.6	874	6	ABQ44403	Abq44403 Oligonuc1
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16	129	11.9	658	12	ADG45405	Adg45405 Liver inf
17	129	11.9	658	12	ADH222707	Adh222707 Partial D
18	127	11.7	579	10	ADH26628	Adh26628 Human Bax
19	127	11.7	579	2	AAZ19764	Aaz19764 Human wil
20	127	11.7	579	4	AAC84598	Aac84598 Human Bax
21	127	11.7	579	6	AAC90811	Aac90811 Human Bax
22	127	11.7	579	6	ABV78149	Abv78149 Human Bax
23	127	11.7	579	6	ABZ35725	Abz35725 Human Bax
24	127	11.7	579	6	ABX09968	Abx09968 Human Bax
25	127	11.7	579	6	ABL91690	AbL91690 Human pol
26	127	11.7	579	9	ADA20845	Ada20845 Human BAX
27	127	11.7	579	9	AAI61054	Aai61054 Human Bax
28	127	11.7	579	11	ADI32061	Adi32061 Human cDN
29	127	11.7	579	12	ADO05979	Ado05979 Pro-apopt
30	127	11.7	624	2	AAQ97606	Aaq97606 Human Bax
31	127	11.7	779	2	AAZ15935	Aaz15935 Human gen
32	127	11.7	791	2	AAZ77545	Aaz77545 Human ova
33	127	11.7	996	2	AAZ06732	Aaz06732 Nucleotid
34	126	11.6	677	9	ADA20844	Ada20844 Human BAX
35	125	11.5	624	2	AAV84005	Aav84005 cDNA enco
36	125	11.5	624	4	AAF77704	Aaf77704 Human bcl
37	120	11.1	596	2	ADF39373	Adf39373 Human BAX
38	120	11.1	1607	2	ADF39391	Adf39391 Human BAX
39	119	11.0	579	10	ADB52764	AdB52764 Primary r
40	119	11.0	579	10	ABT41943	Abt41943 Toxicity
41	118	10.9	579	6	AAF88666	Aaf88666 C. albica
42	118	10.9	583	2	AAT96577	Aat96577 Human bcl
43	118	10.9	583	2	AAZ51134	Aaz51134 Human bcl
44	117.5	10.8	581	2	AAT96578	Aat96578 Mouse bcl
45	117.5	10.8	581	2	AAZ51335	Aaz51335 Mouse bcl

ALIGNMENTS

RESULT 1
AAD46683
ID AAD46683 standard; DNA; 887 BP.
XX
AC AAD46683;
XX
DT 27-JAN-2003 (first entry)
XX
DE Human Bcl-B DNA.
XX
KW Human; Bcl-2; Bcl-B; therapy; apoptosis; cell degenerative disorder;
KW proliferative disorder; muscle degeneration; Alzheimer's disease; CJD;
KW Creutzfeldt-Jacob's disease; Machado-Joseph disease; MJD; transgenic;
KW Parkinson's disease; Huntington's disease; HD; spinocerebellar ataxia;
KW SCA; dentatorubropallidoluysian atrophy; DRPLA; Kennedy's disease;
KW stroke; ischaemia; head trauma; neuroprotection; cytostatic; immunosuppressive; vasotropic;
KW cerebroprotective; autoimmune disorder; chromosome 15; gene; ds.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT CDS 50..864
FT /tag= a
FT /product= "Human Bcl-B protein"

CC activity (e.g. wound healing and tissue repair, ulcers, burns,
CC periodontal disease); activin/inhibin activity; chemotactic/chemokinetic
CC activity; haemostatic and thrombolytic activity (e.g. treating
CC haemophilias); receptor/ligand activity; anti-inflammatory activity; and
CC tumour inhibition activity. The polynucleotides are also stated to be
CC useful for gene therapy. Other activities include inhibiting infections
CC caused by bacteria, fungi, viruses and other parasites (e.g. Hepatitis,
CC malaria); effecting bodily characteristics such as, e.g. weight, colour,
CC skin, effecting biorhythms or cardiac cycles; enhancing fertility;
CC treatment of depression; treatment of pain; hormonal or endocrine
CC activity. The polynucleotides may also be used for recombinant expression
CC of the protein
XX
SQ Sequence 1168 BP; 276 A; 312 C; 300 G; 280 T; 0 U; 0 Other;

Alignment Scores:
Pred. No.: 1,09e-92 Length: 1168
Score: 1045.00 Matches: 136
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 96.40% Indels: 0
DB: 3 Gaps: 0

US-10-071-174-2 (1-204) x AA290049 (1-1168)

QY 9 ThrThrMetAlaAspProLeuArgGluArgThrGluLeuLeuLeuAlaAspTyrLeuGly 28
DB 1 ACCACCATGGCCGACCGCTGGGGAGCGACCGAGCTGTTGTCGCCGACTACCTGGGG 60
QY 29 TyrCysAlaArgGluProGlyThrProGluProAlaProSerThrProGluAlaAlaVal 48
DB 61 TACTGCGCGCGGAACCGCGACCCCGAGCGCGGCCATCCACGCCGAGGCCCGCGGTG 120
QY 49 LeuArgSerAlaAlaAlaArgLeuArgGlnIleHisArgSerPhePheSerAlaTyrLeu 68
DB 121 CTGCGCTCGCGCGCGCCAGGTACGCCAGATTACCGGTCTCTTTTTCGCCCTACCTC 180
QY 69 GlyTyrProGlyValAsnArgPheGluLeuValAlaLeuMetAlaAspSerValLeuSerAsp 88
DB 181 GGCTACCCCGGAGACCGCTTCGAGCTGGTGGCGCTGATGGCGGATTCGCTCTCCGAC 240
QY 89 SerProGlyProThrTrpGlyArgValValThrLeuValThrPheAlaGlyThrLeuLeu 108
DB 241 AGCCCGCGGCCACCTGGGGCAGAGTGGTGACGCTCGTGACCTTCGACGGAGCGCTGTG 300
QY 109 GluArgGlyProLeuValThrAlaArgTyrLysTyrGlyPheGlnProArgLeuLys 128
DB 301 GAGAGAGGCGCGCTGTGTGACCCCGCGGTGGAAGAAGTGGGGCTTCACGCCGCGCTAAAG 360
QY 129 GluGlnGluGlyAspValAlaAlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArg 148
DB 361 GAGCAGAGGGCGCGCTGCGCCCGGAGCTGCCAGCGCTGGTGGCTTGTGAGCTCGCGG 420
QY 149 LeuMetGlyGlnHisArgAlaTrpLeuGlnAlaGlnGlyTyrTrpAspGlyPheCysHis 168
DB 421 CTCATGGGCGACGACCGCGCTGGCTGCGAGGCTCAGGGCGGCTGGGATGGCTTTGTCTAC 480
QY 169 PhePheArgThrProPheProLeuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeu 188
DB 481 TTCTTCAGACCCCTTCCATCGCTTTTGGAGAAACAGCTGGTCAGGCTTTTCTG 540
QY 189 SerCysLeuLeuThrThAlaPheLeuTyrLeuTyrThrArgLeuLeu 204
DB 541 TCATGCTTGTAAACACAGCGCTTATTTATCTCTCGACACGATTATTA 588

RESULT 3
AA290039
ID AA290039 standard; cDNA; 582 BP.
XX
AC AA290039;
XX
DT 09-MAY-2000 (first entry)
XX

DE
XX
KW Hydrophobic domain containing protein clone HP02403 coding sequence.
KW Hydrophobic domain; clone HP02403; nutritional supplement; SCID; HIV;
KW cell proliferation; immune stimulant; immune deficiency; tumour; pain;
KW rheumatoid arthritis; insulin dependent diabetes mellitus; fertility;
KW myasthenia gravis; haematopoiesis regulator; tissue growth; depression;
KW anti-inflammatory; infection; bodily characteristic; ss.
OS Homo sapiens.
XX
PN WO200000506-A2.
XX
PD 06-JAN-2000.
XX
PF 18-JUN-1999; 99WO-JP003242.
PR 26-JUN-1998; 98JP-00180008.
XX
PA (SAGA) SAGAMI CHEM RES CENT.
XX (PROT-) PROTEGENE INC.
PI Kato S, Kimura T;
XX
DR WPI; 2000-160665/14.
DR P-PSDB; AAY78802.
PT
PT Novel human proteins having hydrophobic domains used for research and
diagnostic purposes.
PS
XX Claim 3; Page 84; 117pp; English.

This sequence represents the hydrophobic domain containing protein, clone
HP00631 coding region. The sequence is isolated from a human stomach
cancer cell line. The HP02403 protein contains one putative transmembrane
domain. The protein shows homology to the Japanese quail apoptosis
regulator NR-13. The invention relates to human proteins with hydrophobic
domains, the DNA and the cDNA encoding them. The polynucleotides and
proteins are predicted to have biological activities which make them
suitable for treating, preventing or ameliorating medical conditions in
humans and animals. Suggested activities include nutritional activity
(nutritional source or supplement); cytokine and cell
proliferation/differentiation activity; immune stimulating (e.g. as
vaccines) or suppressing activity (e.g. to treat various immune
deficiencies such as SCIDS or HIV, connective tissue disease, systemic
lupus erythematosus, rheumatoid arthritis, autoimmune pulmonary
inflammation, Guillain-Barre syndrome, myasthenia gravis, graft-versus-host disease
and autoimmune inflammatory eye disease, as well as asthma, allergies and
organ transplantation); haematopoiesis regulating activity (e.g. in
treatment of myeloid or lymphoid cell deficiencies); tissue growth
activity (e.g. wound healing and tissue repair, ulcers, burns,
periodontal disease); activin/inhibin activity; chemotactic/chemokinetic
activity; haemostatic and thrombolytic activity (e.g. treating
haemophilias); receptor/ligand activity; anti-inflammatory activity; and
tumour inhibition activity. The polynucleotides are also stated to be
useful for gene therapy. Other activities include inhibiting infections
caused by bacteria, fungi, viruses and other parasites (e.g. Hepatitis,
malaria); effecting bodily characteristics such as, e.g. weight, colour,
skin, effecting biorhythms or cardiac cycles; enhancing fertility;
treatment of depression; treatment of pain; hormonal or endocrine
activity. The polynucleotides may also be used for recombinant expression
of the protein

SQ Sequence 582 BP; 85 A; 193 C; 191 G; 113 T; 0 U; 0 Other;
Alignment Scores:
Pred. No.: 4.28e-92 Length: 582
Score: 1035.00 Matches: 194
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 95.48% Indels: 0
DB: 3 Gaps: 0

US-10-071-174-2 (1-204) x AA920039 (1-582)	PA	(HUMA-) HUMAN GENOME SCI INC.
QY 11 MetAlaAspProLeuArgGluArgThrGluLeuLeuLeuAlaAspTyrLeuGlyTyrCys 30	XX	Ruben SM, Duan DR, Ni J;
Db 1 ATGGCCGACCGCTGGGAGCGCACCGAGCTGTTCCTGGCCGACTACCTGGGCTACTGC 60	XX	WPI; 2001-476279/51.
QY 31 AlaArgGluProGlyThrProGluProAlaProSerThrProGluAlaAlaValLeuArg 50	DR	P-PSDB; AAB85666.
Db 61 GCCCGGAACCGCGACCCCGGCGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGG 120	XX	Nucleic acids encoding human Bcl-2-like polypeptides, useful for preventing, diagnosing and/or treating.
QY 51 SerAlaAlaAlaArgLeuArgGlnIleHisArgSerPheSerAlaTyrLeuGlyTyr 70	XX	Claim 1; Page 276; 285pp; English.
Db 121 TCGCGCGCGCCGAGTTACGGCAGATTACCGGCTCTTTCTCGCGCTACCTCGGCTAC 180	XX	The invention provides nucleic acid molecules (NAM1) encoding 4 human Bcl-2-like polypeptides (PEP1). The NAM1 and PEP1 may be used in the prevention, diagnosis and treatment of diseases associated with inappropriate Bcl-2-like polypeptides, expression. The NAM1 may be used to produce the soluble Bcl-2-like polypeptides by standard recombinant methodology. The polypeptides may also be used as antigens in the production of antibodies against Bcl-2 and in assays to identify modulators of Bcl-2 expression and activity. The anti-Bcl-2 antibodies and antagonists may be used to down regulate expression and activity. The anti-PEP1 antibodies may also be used as diagnostic agents for detecting the presence of Bcl-2 polyPs in samples (e.g. by enzyme linked immunosorbant assay (ELISA)). Disorders that may be prevented, diagnosed and/or treated by the above methods include, immunodeficiencies (e.g. a gammaglobulinemia and B cell lymphoproliferative disorder), allergic disorders (e.g. rheumatoid arthritis and Grave's disease), allergic reactions, inflammations, respiratory diseases and cardiovascular disorders (a full list of disorders is given in the specification). The present sequence represents a human Bcl-2-like polypeptide encoding cDNA
QY 91 GlyProThrTyrGlyArgValValThrLeuValThrPheAlaGlyThrLeuLeuGluArg 110	CC	Sequence 522 BP; 102 A; 148 C; 154 G; 118 T; 0 U; 0 Other;
Db 241 GCGCCACCTGGGCGAGAGTGCTGACCTCGTACCTTCGAGGAGCGCTGCTGAGAGA 300	XX	Alignment Scores:
QY 111 GlyProLeuValThrAlaArgTyrLysLysTyrGlyPheGlnProArgLeuLysGluGln 130	XX	Pred. No.: 1.96e-59
Db 301 GGGCCGCTGTGACCGCGGTGGAGAAAGTGGGGCTTCAGCCGCGGCTAAAGGAGCAG 360	XX	Length: 522
QY 131 GluGlyAspValAlaArgAspCysGlnArgLeuValAlaLeuSerSerArgLeuMet 150	XX	Matches: 130
Db 361 GAGGGGACGTCGCCGGGACTCCAGCGCTGGTGGCTTGTGAGCTCGCGGCTCATG 420	XX	Score: 701.00
QY 151 GlyGlnHisArgAlaTrpLeuGlnAlaGlnGlyGlyTrpAspGlyPheCysHisPhe 170	XX	Percent Similarity: 100.00%
Db 421 GGGCAGCACCGCTGGCTGCGAGGCTCAGGGCGGCTGGGCTTTGTCACTCTCTC 480	XX	Best Local Similarity: 100.00%
QY 171 ArgThrProPheProLeuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeuSerCys 190	XX	Query Match: 64.67%
Db 481 AGACCCCTTTTCCACGCGCTTTTGGAGAAACAGCTGCTGTCAGGCTTTTCTGTATGC 540	XX	Indels: 0
QY 191 LeuLeuThrThrAlaPheIleTyrLeuTrpThrArgLeuLeu 204	XX	Gaps: 4
Db 541 TTGTTACACAGCTTTTCACTTATCTCTGGACACGATTATTA 582	XX	US-10-071-174-2 (1-204) x AAH47022 (1-522)
RESULT 4	QY	75 PheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerProGlyProThrTrp 94
AAH47022	Db	3 TTCGAGCTGGTGGCGCTGATGGGGATTCCGTGCTCTCCGACAGCCCGCGCCACCTGG 62
AC	QY	95 GlyArgValValThrLeuValThrPheAlaGlyThrLeuLeuGluArgGlyProLeuVal 114
XX	Db	63 GGCAGAGTGGTGGCGCTCGTACCTTCGAGGGGACGCTGCTGGAGAGGGCGGCTGGTG 122
DT	QY	115 ThrAlaArgTyrLysLysTyrGlyPheGlnProArgLeuLysGluGlnGlyAspVal 134
XX	Db	123 ACCGCCCGGTGGAGAGTGGGGCTTCAGCCGCGCTAAAGAGCAGGAGGGCGACGTC 182
DE	QY	135 AlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArgLeuMetGlyGlnHisArg 154
XX	Db	183 GCGCGGAGCTGCCAGCGCTGGTGGCTTGTGAGCTCGCGGCTCATGGGGCAGCACCGC 242
XX	QY	155 AlaTrpLeuGlnAlaGlnGlyTrpAspGlyPheCysHisPhePheArgThrProPhe 174
XX	Db	243 GCTTGGCTGCGAGGCTCAGGGCGGCTGGGATGGCTTTGTGCTCTTTCAGGACCCCTTT 302
XX	QY	175 ProLeuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeuSerCysLeuLeuThrThr 194
XX	Db	303 CCACCTGGCTTTTGGAGAAAACAGCTGCTCCAGGCTTTTCTGTGTCATGCTTTTAAACA 362
XX	QY	195 AlaPheIleTyrLeuTrpThrArgLeuLeu 204
XX	Db	363 GCTTTCATTATCTCTGGACACGATTATTA 392
XX	XX	RESULT 5
XX	XX	ABK41913

ID ABK41913 standard; cDNA; 548 BP.
XX
AC ABK41913;
XX
DT 21-MAY-2002 (first entry)
XX
DE cDNA encoding novel human connective tissue related polypeptide #301.
XX
KW Human; connective tissue related disorder; cancer; gene therapy;
KW cytotstatic; gene; ss.
XX
OS Homo sapiens.
XX
PN WO20015343-A1.
XX
PD 02-AUG-2001.
XX
PF 17-JAN-2001; 2001WO-US001322.
XX
PP 31-JAN-2000; 2000US-0179065P.
XX
PR 04-FEB-2000; 2000US-0180628P.
PR 24-FEB-2000; 2000US-0184664P.
PR 02-MAR-2000; 2000US-0186350P.
PR 16-MAR-2000; 2000US-0189874P.
PR 17-MAR-2000; 2000US-0190076P.
PR 18-APR-2000; 2000US-0198123P.
PR 19-MAY-2000; 2000US-0205515P.
PR 07-JUN-2000; 2000US-0209467P.
PR 28-JUN-2000; 2000US-0214886P.
PR 30-JUN-2000; 2000US-0215135P.
PR 07-JUL-2000; 2000US-0216647P.
PR 07-JUL-2000; 2000US-0216880P.
PR 11-JUL-2000; 2000US-0217487P.
PR 11-JUL-2000; 2000US-0217496P.
PR 14-JUL-2000; 2000US-0218290P.
PR 26-JUL-2000; 2000US-0220963P.
PR 26-JUL-2000; 2000US-0220964P.
PR 14-AUG-2000; 2000US-0224518P.
PR 14-AUG-2000; 2000US-0224519P.
PR 14-AUG-2000; 2000US-0225213P.
PR 14-AUG-2000; 2000US-0225214P.
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PR 14-AUG-2000; 2000US-0225447P.
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PR 18-AUG-2000; 2000US-0226279P.
PR 22-AUG-2000; 2000US-0226681P.
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PR 30-AUG-2000; 2000US-0228924P.
PR 01-SEP-2000; 2000US-0229287P.
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PR 05-SEP-2000; 2000US-0229509P.
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PR 06-SEP-2000; 2000US-0230433P.
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PR 08-SEP-2000; 2000US-0233088P.
PR 08-SEP-2000; 2000US-0233089P.
PR 12-SEP-2000; 2000US-0231968P.
PR 14-SEP-2000; 2000US-0232397P.
PR 14-SEP-2000; 2000US-0232398P.
PR 14-SEP-2000; 2000US-0232399P.
PR 14-SEP-2000; 2000US-0232400P.
PR 14-SEP-2000; 2000US-0232401P.
PR 14-SEP-2000; 2000US-0233063P.
PR 14-SEP-2000; 2000US-0233064P.
PR 14-SEP-2000; 2000US-0233065P.
PR 21-SEP-2000; 2000US-0234223P.
PR 21-SEP-2000; 2000US-0234274P.
PR 25-SEP-2000; 2000US-0234997P.
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PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
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PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
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PR 13-OCT-2000; 2000US-0239935P.
PR 13-OCT-2000; 2000US-0239937P.
PR 20-OCT-2000; 2000US-0240560P.
PR 20-OCT-2000; 2000US-0241221P.
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PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 20-OCT-2000; 2000US-0241826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
PR 08-NOV-2000; 2000US-0246476P.
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PR 08-NOV-2000; 2000US-0246478P.
PR 08-NOV-2000; 2000US-0246523P.
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PR 08-NOV-2000; 2000US-0246609P.
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PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
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PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
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PR 17-NOV-2000; 2000US-0249246P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249299P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.

05-DEC-2000; 2000US-0256719P.
06-DEC-2000; 2000US-0251479P.
08-DEC-2000; 2000US-0251856P.
08-DEC-2000; 2000US-0251868P.
08-DEC-2000; 2000US-0251869P.
08-DEC-2000; 2000US-0251989P.
08-DEC-2000; 2000US-0251990P.
11-DEC-2000; 2000US-0254097P.
05-JAN-2001; 2001US-0259678P.
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Rosen CA, Barash SC, Ruben SM;
XX WPI; 2001-565190/63.
XX P-PSDB; AAU86735.
XX Nucleic acid encoding novel connective tissue associated polypeptides,
XX used in diagnosing, preventing, treating or ameliorating a disorder such
XX as cancer or rheumatoid arthritis.
XX
XX Claim 4; SEQ ID NO 311; 673pp; English.
XX
XX The present invention relates to the isolation of novel human connective
XX tissue related polypeptides (AAU86435-AAU86923) and the polynucleotide
XX (cDNA and genomic) sequences encoding them. The sequences of the
XX invention are useful in the diagnosis, treatment, prevention and/or
XX prognosis of diseases associated with connective tissue(s), including
XX cancer. The polynucleotide sequences of the invention are also useful in
XX gene therapy. ABK41613-ABK42101 represent cDNA sequences encoding the
XX novel human connective tissue related polypeptides. Note: The sequence
XX data for this patent did not form part of the printed specification, but
XX was obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX
XX Sequence 548 BP; 99 A; 154 C; 167 G; 121 T; 0 U; 7 Other;

Alignment Scores:
Pred. No.: 2,47e-53 Length: 548
Score: 639.00 Matches: 127
Percent Similarity: 97.69% Conservative: 0
Best Local Similarity: 97.69% Mismatches: 3
Query Match: 58.95% Indels: 2
DB: 4 Gaps: 0

US-10-071-174-2 (1-204) x ABK41913 (1-548)

QY 75 PheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerProGlyProThrTrp 94
DB 74 TTGAGCTGGTGGCTGATGGGGGATTCGCTCTCCGACAGCCCGCCACCTGG 133
QY 95 GlyArgValValThrLeuValThrPheAlaGlyThrLeuLeuGluArgGlyProLeuVal 114
DB 134 GC-AGAGTGGTGGCTCGGACCTTCGCGAGGACGCTGCTGGAGAGAGCGCGCTGGTG 192
QY 115 ThrAlaArgTrpLysTrpGlyPheGlnProArgLeuLysGluGlnGlyAspVal 134
DB 193 ACCCGCGGTGGAGAGAGTGGGCTTCCAGCCGGCTTAAAGGAGCAGGAGGCGCAGTC 252
QY 135 AlaArgAspCysGlnArgLeuValAlaLeuSerSerArgLeuMetGlyGlnHisArg 154
DB 253 GCSSGGACCTGCCAGCGCTTGGTGGCTTGTGAGCTCGCGCTCATGGGCGACACCGG 312
QY 155 AlaTrpLeuGlnAlaGlnGlyTrpAspGlyPheCysHisPheArgThrProPhe 174
DB 313 CC-TGGCTGCARGCTCAGGCGGCTGGATGGCTTTTGTCACTTCTTCAGGACCCCTTT 371
QY 175 ProteuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeuSerCysLeuLeuThrThr 194
DB 372 CCACGTGCTTTTGGAGAAAACAGCTGGTCCAGGCTTTTGTGTCATGCTTGTTAACACA 431
QY 195 AlaPheIleTyLeuTrpThrArgLeuLeu 204

Db 432 GCCTTCATTATCTCTGGACACGATTATTA 461
RESULT 6
ADBS9580
ID ADBS9580 standard; cDNA; 548 BP.
XX AC ADBS9580;
XX
DT 04-DEC-2003 (first entry)
XX
DE Connective tissue related polynucleotide #301.
XX
XX cytostatic; neuroprotective; nootropic; antiparkinsonian; cardiovascular;
XX antiarteriosclerotic; immunosuppressive; antirheumatic; antiarthritic;
XX antiinflammatory; antiallergic; antiasthmatic; dermatological;
XX nephrotropic; virucide; fungicide; antibacterial; antiparasitic;
XX gene therapy; ds; connective tissues disorder; rheumatoid arthritis;
XX systemic lupus erythematosus; scleroderma; Sjogren's syndrome; cancer;
XX cancer metastasis; neoplasia; leukaemia; neurodegenerative disorder;
XX Alzheimer's disease; Parkinson's disease; cardiovascular disease;
XX atherosclerosis; myocarditis; cardiopulmonary bypass complication;
XX autoimmune disease; multiple sclerosis; allergic reaction; asthma;
XX rhinitis; eczema; inflammatory condition; Crohn's disease; nephritis;
XX gastrointestinal disorder; inflammatory bowel disease;
XX organ transplant rejection; immune system disorder; Bruton's disease;
XX X-linked lymphoproliferative syndrome;
XX B-cell lymphoproliferative disorder; HIV; AIDS; infection;
XX chromosome identification; chromosome mapping;
XX connective tissue related polynucleotide; gene; as.
XX
OS Homo sapiens.
XX
XX US2003054375-A1.
XX
XX 20-MAR-2003.
XX
XX 07-MAR-2002; 2002US-00092154.
XX
XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184664P.
XX 02-MAR-2000; 2000US-0186350P.
XX 16-MAR-2000; 2000US-0189874P.
XX 17-MAR-2000; 2000US-0190076P.
XX 18-APR-2000; 2000US-0198123P.
XX 19-MAY-2000; 2000US-0205515P.
XX 07-JUN-2000; 2000US-0209467P.
XX 30-JUN-2000; 2000US-0215135P.
XX 07-JUL-2000; 2000US-0216647P.
XX 07-JUL-2000; 2000US-0216880P.
XX 11-JUL-2000; 2000US-0217487P.
XX 11-JUL-2000; 2000US-0217496P.
XX 14-JUL-2000; 2000US-0218290P.
XX 26-JUL-2000; 2000US-0220963P.
XX 26-JUL-2000; 2000US-0220964P.
XX 14-AUG-2000; 2000US-0224518P.
XX 14-AUG-2000; 2000US-0224519P.
XX 14-AUG-2000; 2000US-0225213P.
XX 14-AUG-2000; 2000US-0225214P.
XX 14-AUG-2000; 2000US-0225266P.
XX 14-AUG-2000; 2000US-0225267P.
XX 14-AUG-2000; 2000US-0225268P.
XX 14-AUG-2000; 2000US-0225270P.
XX 14-AUG-2000; 2000US-0225447P.
XX 14-AUG-2000; 2000US-0225757P.
XX 14-AUG-2000; 2000US-0225758P.
XX 14-AUG-2000; 2000US-0225759P.
XX 18-AUG-2000; 2000US-0226279P.
XX 22-AUG-2000; 2000US-0226681P.
XX 22-AUG-2000; 2000US-0226868P.
XX 22-AUG-2000; 2000US-0227182P.
XX 23-AUG-2000; 2000US-0227009P.

PR 30-AUG-2000; 2000US-0228924P.
PR 01-SEP-2000; 2000US-0229287P.
PR 01-SEP-2000; 2000US-0229343P.
PR 01-SEP-2000; 2000US-0229344P.
PR 01-SEP-2000; 2000US-0229345P.
PR 01-SEP-2000; 2000US-0229346P.
PR 05-SEP-2000; 2000US-0229513P.
PR 05-SEP-2000; 2000US-0230437P.
PR 06-SEP-2000; 2000US-0230438P.
PR 08-SEP-2000; 2000US-0231242P.
PR 08-SEP-2000; 2000US-0231243P.
PR 08-SEP-2000; 2000US-0231244P.
PR 08-SEP-2000; 2000US-0231413P.
PR 08-SEP-2000; 2000US-0231414P.
PR 08-SEP-2000; 2000US-0232080P.
PR 08-SEP-2000; 2000US-0232081P.
PR 12-SEP-2000; 2000US-0231968P.
PR 14-SEP-2000; 2000US-0232397P.
PR 14-SEP-2000; 2000US-0232398P.
PR 14-SEP-2000; 2000US-0232399P.
PR 14-SEP-2000; 2000US-0232400P.
PR 14-SEP-2000; 2000US-0232401P.
PR 14-SEP-2000; 2000US-0233063P.
PR 14-SEP-2000; 2000US-0233064P.
PR 14-SEP-2000; 2000US-0233065P.
PR 21-SEP-2000; 2000US-0234223P.
PR 21-SEP-2000; 2000US-0234274P.
PR 25-SEP-2000; 2000US-0234597P.
PR 25-SEP-2000; 2000US-0234598P.
PR 26-SEP-2000; 2000US-0235484P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 02-OCT-2000; 2000US-0237040P.
PR 13-OCT-2000; 2000US-0239335P.
PR 13-OCT-2000; 2000US-0239337P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241786P.
PR 20-OCT-2000; 2000US-0241787P.
PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 01-NOV-2000; 2000US-0241826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
PR 08-NOV-2000; 2000US-0246476P.
PR 08-NOV-2000; 2000US-0246477P.
PR 08-NOV-2000; 2000US-0246478P.
PR 08-NOV-2000; 2000US-0246523P.
PR 08-NOV-2000; 2000US-0246524P.
PR 08-NOV-2000; 2000US-0246525P.
PR 08-NOV-2000; 2000US-0246526P.
PR 08-NOV-2000; 2000US-0246527P.
PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246532P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
PR 17-NOV-2000; 2000US-0249210P.

PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.
PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
PR 17-NOV-2000; 2000US-0249245P.
PR 17-NOV-2000; 2000US-0249246P.
PR 17-NOV-2000; 2000US-0249255P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249299P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251868P.
PR 08-DEC-2000; 2000US-0251869P.
PR 08-DEC-2000; 2000US-0251989P.
PR 08-DEC-2000; 2000US-0251990P.
PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.
PR 17-JAN-2001; 2001US-00764847.

XX (HUNA-) HUMAN GENOME SCI INC.

PA Rosen CA, Ruben SM, Barash SC;

XX WPI; 2003-634869/60.

DR P-FSDB; ADB60069.

PT New connective tissue-related polypeptides and polynucleotides, useful
PT for treating, preventing and/or prognosing e.g. disorders of connective
PT tissue, (e.g. rheumatoid arthritis), cancers, cancer metastases and/or
PT neoplasias.

XX Claim 1; SEQ ID NO 311; 248pp; English.

XX The invention describes an isolated nucleic acid molecule (I), which
CC comprises a sequence that is at least 95 % identical to a connective
CC tissue-related polynucleotide encoding connective tissue antigens (CTA).
CC The polypeptide or polynucleotide is useful for preventing, treating, or
CC ameliorating medical conditions in a mammal. The connective tissue
CC polypeptides, polynucleotides and antibodies are particularly useful for
CC treating, preventing and/or prognosing disorders of connective tissues
CC (e.g. rheumatoid arthritis, discoid and systemic lupus erythematosus,
CC scleroderma, or Sjogren's syndrome), cancers, cancer metastases and/or
CC neoplasias (e.g. leukaemia), neurodegenerative disorders (e.g.
CC Alzheimer's disease, or Parkinson's disease), cardiovascular diseases
CC (e.g. atherosclerosis, myocarditis or cardiopulmonary bypass
CC complications), autoimmune diseases (e.g. systemic lupus erythematosus,
CC rheumatoid arthritis, or multiple sclerosis), allergic reactions (e.g.

Alignment Scores:

Pred. No.:	2,47e-53	Length:	548
Score:	639.00	Matches:	127
Percent Similarity:	97.69%	Conservative:	0
Best Local Similarity:	97.69%	Mismatches:	3
Query Match:	58.95%	Indels:	2
DB:	9	Gaps:	0

US-10-071-174-2 (1-204) x ADB59580 (1-548)

QY 75 PhecluleuValAlaLeuMetAlaAspSerValIleuSerAspSerProGlyProThrTyr 94

DB 74 TTCGAGCTGGTGGCGCTGA*GGCGGATTCGCTCTCCGACAGCCCGCCACCTGG 133

QY	95	GlyArgValValThrLeuValThrPheAlaGlyThrIleuLeuGluArgGlyProIleuVal	114
DB	134	GC-AGAGTGGTGCAGCGTCGTGACCTTCGCAGGACGCTGCTGGAGAGAGGGCGCTGGTG	192
QY	115	ThrAlaArgTTPLeuLysLysTTPGlyPheGlnProArgIleuLysGlnGlnGlyAspVal	134
DB	193	ACCGCCCGGTGGAGAGTGGGGCTCCACCGCGGCTAAGGAGCAGGAGGCGCACGTC	252
QY	135	AlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArgLeuMetGlyGlnHisArg	154
DB	253	GCCSGGACTCCACAGCGCTGTGTCCTTCTGCTGAGCTCGCGGCTCATGGGCGACACCGG	312
QY	155	AlaTTPLeuGlnAlaGlnGlyGlyTTPAspGlyPheCysHisPhePheArgThrProPhe	174
DB	313	CC-TGGCTGCARGCTCAGGGCGGTGGGATGGCTTTTGTCACTTCTTCAGGACCCCCCTT	371
QY	175	ProLeuAlaPheTTPArgLysGlnLeuValGlnAlaPheLeuSerCysLeuLeuThrThr	194
DB	372	CCACTGGCTTTTGGAGAAACACACTGGTCCAGGCTTTTCTGTCACTGCTTGTAAACACA	431
QY	195	AlaPheIleTyrLeuTTPThrArgLeuLeu	204
DB	432	GCCTTCATTATCTCTGGACACGATTATTA	461
RESULT 7			
ABQ44403			
ID ABQ44403 standard; DNA; 874 BP.			
XX	AC		
XX	ABQ44403;		
XX	DT		
XX	12-JUL-2002 (first entry)		
XX	DE		
XX	Oligonucleotide for detecting cytosine methylation SEQ ID NO 30994.		
XX	Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;		
KW	drug; side effect; cancer; central nervous system; cardiovascular;		
KW	gastrointestinal; respiratory system; single nucleotide polymorphism;		
KW	SNP; cell differentiation; ds.		
XX	Homo sapiens.		
OS	WO200218632-A2.		
XX	PN		
XX	07-MAR-2002.		
XX	PF		
XX	01-SEP-2001; 2001WO-EP010074.		
XX	PR		
XX	01-SEP-2000; 2000DE-01043826.		
XX	05-SEP-2000; 2000DE-0104543.		
XX	XX		
PA	(EPIG-) EPIGENOMICS AG.		
XX	PI		
XX	olek A, Piepenbrock C, Berlin K, Guetig D;		
XX	WPI; 2002-371829/40.		

XX Determining the degree of cytosine methylation in genomic DNA, useful for
 PT diagnosis and prognosis, comprises selective hybridization of amplicons
 PT from chemically treated DNA.
 XX
 XX Claim 12; 56pp + Sequence Listing; 56pp; German.
 PS
 XX This invention describes a novel method for determining the degree of
 CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 CC genomic sample of DNA. The sample is treated chemically to convert
 CC cytosine (C) but not methylated C, to uracil, then part of the genomic
 CC DNA that contains the target C is amplified to form a labeled amplicon.
 CC The amplicon is hybridized to two classes, each with at least one member,
 CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
 CC degree of hybridisation to both classes is determined from the label on
 CC the amplicon. From the ratio of labels hybridised to the two classes of
 CC oligomers, the degree of methylation is calculated. The method is used:
 CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs

	DB	678	GTAACCTTCGCAAAACCGCTACTAAAAAACAACGCCTAAATAACGCCGTATAAAAAGC	367
PB	XX	FF		
PF	XX			
PP	XX			
PR	XX			
PS	XX			
PT	XX			
(EPIG-)	PA	EPIGENOMICS AG.		
Olek A,	Db	Piepenbrock C, Berlin K, Guetig D;		
WPI; 2002-371829/40.	DR			
Determining the degree of cytosine methylation in genomic DNA, useful for diagnosis and prognosis, comprises selective hybridization of amplicons from chemically treated DNA.	CC			
Claim 12; 56pp + Sequence Listing; 56pp; German.	CC			
This invention describes a novel method for determining the degree of methylation of a particular cytosine in a motif 5'- CpG -3', present in a genomic sample of DNA. The sample is treated chemically to convert cytosine (C) but not methylated C, to uracil, then part of the genomic DNA that contains the target C is amplified to form a labeled amplicon. The amplicon is hybridised to two classes, each with at least one member, of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the degree of hybridisation to both classes is determined from the label on the amplicon. From the ratio of labels hybridized to the two classes of oligomers, the degree of methylation is calculated. The method is used:	CC			
(i) for diagnosis and/or prognosis of side effects of therapeutic drugs and of a wide range of diseases, e.g. cancer, disorders of the central nervous system, cardiovascular, gastrointestinal and respiratory systems etc., particularly by detecting mutations or single nucleotide polymorphisms (SNPs); and (ii) for differentiation of cell or tissue types and for investigating cell differentiation. The method allows the methylation status of many C residues to be determined simultaneously. ABQ13410-	CC			
ABQ54121 represent genomic DNA sequences used to illustrate the method for determining the degree of cytosine methylation described in the disclosure of the invention	CC			
SQ	Sequence 874 BP; 119 A; 90 C; 291 G; 374 T; 0 U; 0 Other;			
Alignment Scores:				
Pred. No.: 3.09e-45 Length: 874				
Score: 559.00 Matches: 110				
Percent Similarity: 83.75% Conservative: 24				
Best Local Similarity: 68.75% Mismatches: 26				
Query Match: 51.57% Indels: 0				
DB: 6 Gaps: 0				
US-10-071-174-2 (1-204) x ABQ44402 (1-874)				
QY	1 MetValAspGlnLeuArgGluThrMetAlaAspProLeuArgGluArgThrGlu 20			
Db	726 ATAATTACCATTACGAAGAAGCACCATACACGGACCGTACGAAACGCCGAA 667			
QY	21 IeuLeuLeuAlaAspTyrlleuglyTyrCyslaarggluProglylthrProgluProala 40			
Db	666 CTATTACTACCGACTACTCTAAATACTACGCCCGCAAACCGCACCCCACCGACG 607			
QY	41 ProSerThrProgluaLaalavalLeuArgGerAlaalaaLagLeuArgGlinileHis 60			
Db	606 CCATCCAGCCCCGAACCGCGTGACTACTGCTCCGGACCGCCAATTTACGNCAANTTCAC 547			
QY	61 ArgSerPheSeraItyrlleuglyTyrProglyAsnArghpeGluLeuValaleu 80			
Db	546 CGATCCTCTTTTTCCGCGCTACCTCGACTACCCCGAAAAACCGCTTCGAACATAATACGCTA 487			
QY	81 MetAlaAspSerValIeuSerAspSerProglyProThrTrpGlyArgValValThrLeu 100			
Db	486 ATAAACGAATTCGTTACTCTCGACACACCCCGACCCACCTAAAAACAAATAATATACGCTC 427			
QY	101 ValThrPheAlaGlYThrLeuLeuGluArgGlyProLeuValThrAlaAgTrpLylys 120			

Qy 101 ValThrPheAlaGlyThrLeuLeuGluArgGlyProLeuValThrAlaArgTrpLysLys 120

Pred. No.: 9,92e-41 Length: 874
 Score: 513.00 Matches: 111
 Percent Similarity: 69.33% Conservative: 2
 Best Local Similarity: 68.10% Mismatches: 50
 Query Match: 47.32% Indels: 0
 DB: 6 Gaps: 0

US-10-071-174-2 (1-204) x ABQ44405 (1-874)

Qy 1 MetValAspGlnLeuArgGluArgThrMetAlaAspProLeuArgGluArgThrGlu 20
 Db 149 ATGTTGATAGTTCGGGAGCGATATTATATGTCGATTCGTCGGGAGCGTATCGAG 208
 Qy 21 LeuLeuLeuAlaAspTyrLeuGlyTyrCysAlaArgGluProGlyThrProGluProAla 40
 Db 209 TTGTTGTGTCGATATTTCGGGTATTCGGTTCGGGAAATTCGATATTCGAGTCGGCG 268
 Qy 41 ProSerThrProGluAlaAlaValLeuArgSerAlaAlaAlaArgLeuArgGlnIleHis 60
 Db 269 TTATTTACGTTTCAGGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCG 328
 Qy 61 ArgSerPhePheSerAlaTyrLeuGlyTyrProGlyAsnArgPheGluLeuValAlaLeu 80
 Db 329 CGGTTTTTTTTCGTTATTTTCGGTTATTCGGGAATCGTTTCGAGTTGTCGGCTG 388
 Qy 81 MetAlaAspSerValLeuSerAspSerProGlyProThrTrpGlyArgValValThrLeu 100
 Db 389 ATGCGGATTTTCGTTTTCGATAGTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTT 448
 Qy 101 ValThrPheAlaGlyThrLeuLeuGluArgGlyProLeuValThrAlaArgTrpIleHis 120
 Db 449 GTGATTTTCGTCGAGCGTTTCGTCGAGAGGTCGTCGTCGTCGTCGTCGTCGTCG 508
 Qy 121 TrpGlyPheGlnProArgLeuLysGluGlnGluGlyAspValAlaArgAspCysGlnArg 140
 Db 509 TGGGTTTTTTCGCGGTTAAAGAGTAGAGGCGGCGAGTCGTCGTCGTCGTCGTCGTCG 568
 Qy 141 LeuValAlaLeuLeuSerSerArgLeuMetGlyGlnHisArgAlaTrpLeuGlnAlaGln 160
 Db 569 TTGTCGTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTT 628
 Qy 161 GlyGlyTrp 163
 Db 629 GCGGTTGG 637

RESULT 10

ABQ44405/c

ID ABQ44405 standard; DNA; 874 BP.

AC ABQ44405;

XX 12-JUL-2002 (first entry)

DT 12-JUL-2002 (first entry)

DE Oligonucleotide for detecting cytosine methylation SEQ ID NO 30996.

XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;

XX drug; side effect; cancer; central nervous system; cardiovascular;

XX gastrointestinal; respiratory system; single nucleotide polymorphism;

XX SNP; cell differentiation; ds.

XX Homo sapiens.

XX WO200218632-A2.

XX 07-MAR-2002.

XX 01-SEP-2001; 2001WO-EP010074..

XX 01-SEP-2000; 2000DE-01043826.

XX 05-SEP-2000; 2000DE-0104543.

XX (EPIG-) EPIGENOMICS AG.

FA

XX

PI Olek A, Piepenbrock C, Berlin K, Guetig D;

XX WPI; 2002-371829/40.

XX Determining the degree of cytosine methylation in genomic DNA, useful for
 diagnosis and prognosis, comprises selective hybridization of amplicons
 from chemically treated DNA.

XX Claim 12; 56pp + Sequence Listing; 56pp; German.

XX This invention describes a novel method for determining the degree of
 methylation of a particular cytosine in a motif 5'-CpG-3', present in a
 genomic sample of DNA. The sample is treated chemically to convert
 cytosine (C) but not methylated C, to uracil, then part of the genomic
 DNA that contains the target C is amplified to form a labeled amplicon.
 The amplicon is hybridised to two classes, each with at least one member,
 of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
 degree of hybridisation to both classes is determined from the label on
 the amplicon. From the ratio of labels hybridised to the two classes of
 oligomers, the degree of methylation is calculated. The method is used:
 (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
 and/or of a wide range of diseases, e.g. cancer, disorders of the central
 nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
 particularly by detecting mutations or single nucleotide polymorphisms
 (SNP's); and (ii) for differentiation of cell or tissue types and for
 investigating cell differentiation. The method allows the methylation
 status of many C residues to be determined simultaneously. ABQ13410-
 ABQ54121 represent genomic DNA sequences used to illustrate the method
 for determining the degree of cytosine methylation described in the
 disclosure of the invention

SQ Sequence 874 BP; 320 A; 320 C; 90 G; 144 T; 0 U; 0 Other;

Alignment Scores:

Pred. No.: 9,92e-41 Length: 874
 Score: 513.00 Matches: 111
 Percent Similarity: 69.33% Conservative: 2
 Best Local Similarity: 68.10% Mismatches: 50
 Query Match: 47.32% Indels: 0
 DB: 6 Gaps: 0

US-10-071-174-2 (1-204) x ABQ44405 (1-874)

Qy 1 MetValAspGlnLeuArgGluArgThrMetAlaAspProLeuArgGluArgThrGlu 20
 Db 726 ATGTTGATAGTTCGGGAGCGATATTATATGTCGATTCGTCGGGAGCGTATCGAG 667
 Qy 21 LeuLeuLeuAlaAspTyrLeuGlyTyrCysAlaArgGluProGlyThrProGluProAla 40
 Db 666 TTGTTGTGTCGATATTTCGGGTATTCGGTTCGGGAATTCGATATTCGAGTCGGCG 607
 Qy 41 ProSerThrProGluAlaAlaValLeuArgSerAlaAlaAlaArgLeuArgGlnIleHis 60
 Db 606 TTATTTACGTTTCAGGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCGTCG 547
 Qy 61 ArgSerPhePheSerAlaTyrLeuGlyTyrProGlyAsnArgPheGluLeuValAlaLeu 80
 Db 546 CGGTTTTTTTTCGTTTATTCGGTTATTCGGGAATTCGTTTCGAGTTGTCGGCTG 487
 Qy 81 MetAlaAspSerValLeuSerAspSerProGlyProThrTrpGlyArgValValThrLeu 100
 Db 486 ATGCGGATTTTCGTTTTCGATAGTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTT 427
 Qy 101 ValThrPheAlaGlyThrLeuLeuGluArgGlyProLeuValThrAlaArgTrpIleHis 120
 Db 426 GTGATTTTCGAGGACGTTTCGAGAGAGGTCGTCGTCGTCGTCGTCGTCGTCGTCG 367
 Qy 121 TrpGlyPheGlnProArgLeuLysGluGlnGluGlyAspValAlaArgAspCysGlnArg 140
 Db 366 TGGGTTTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTTTTCGTT 307
 Qy 141 LeuValAlaLeuLeuSerSerArgLeuMetGlyGlnHisArgAlaTrpLeuGlnAlaGln 160

Db 306 TTGGTGGTTTGTAGTTCGGGTTTANGGGTAGTATCGGTTGGTTGGTTAGGTTAG 247
QY 161 GlyGlyTyr 163
Db 246 GGGGTTGG 238

RESULT 11
ADD34132
ID ADD34132 standard; DNA; 650 BP.
XX
AC ADD34132;
DT 15-JAN-2004 (first entry)
XX
DE Mouse mitochondrial DNA sequence SEQ ID NO:1910.
XX
KW ds; mouse; array; mitochondrial; hybridisation; energy-metabolism;
KW mitochondrial disease; oxidative phosphorylation dysfunction;
KW oxidative stress; apoptosis; aging.
XX
OS Mus musculus.
XX
FN WC2003020220-A2.
XX
PD 13-MAR-2003.
XX
PF 30-AUG-2002; 2002WO-US027886.
XX
PR 30-AUG-2001; 2001US-0316223P.
PR 31-AUG-2001; 2001CA-02356540.
XX
PA (UYEN-) UNIV EMORY.
XX
PI Wallace DC, Levy S, Kerstann K, Procaccio V;
XX WPI; 2003-300821/29.
DR
XX
PT Array containing probes for genes involved in mitochondrial biology,
PT useful for determining mitochondrial biology gene expression profiles for
PT use in diagnosing pathologies and identifying biochemical pathways.
XX
PS Claim 2; SEQ ID NO 1910; 201pp; English.
XX
CC The invention relates to a novel array comprising at least two isolated
CC nucleotide molecules, each molecule having a sequence capable of uniquely
CC hybridising to a nucleic acid molecule which is an expression product of
CC a gene involved in mitochondrial biology. The array comprises two or more
CC isolated nucleic acid molecules or spots, each molecule having a sequence
CC chosen from sequence of 994 human probes and 2046 mouse probes. An array
CC of the invention is useful for determining an expression profile of a
CC mouse or human sample containing nucleic acid, by contacting the array
CC with the sample under conditions allowing selective hybridisation, and
CC measuring hybridisation of nucleic acid in the sample to the array to
CC produce an expression profile. The array is also useful for determining
CC an expression profile of a first labelled sample containing nucleic acid
CC relative to a second, differently labelled sample containing nucleic
CC acid. The second sample is a reference or a standard. An array is useful
CC for determining an expression profile diagnostic of an energy-metabolism-
CC related physiological condition. An array of the invention is useful for
CC determining mitochondrial biology gene expression profiles of organisms,
CC such as human, mice and closely related species, tissue and organs of
CC such organisms, which are useful for determining expression profiles
CC diagnostic of energy metabolism-related physiological conditions,
CC diagnosing such physiological conditions, identifying biochemical
CC pathways, genes, and mutations involved in such physiological conditions,
CC identifying therapeutic agents useful for preventing and/or treating such
CC physiological conditions, evaluating and/or monitoring the efficacy of
CC such therapies, and creating and identifying animal models of human
CC energy metabolism-related physiological conditions. An array is also
CC useful for defining expression signatures or profiles for mitochondrial
CC diseases, as well as distinguishing clinical disorders that result from
CC oxidative phosphorylation (OXPHOS) dysfunction, oxidative stress,
CC apoptosis and aging. An array of the invention contains probes of genes

CC not previously recognised to participate in mitochondrial biology. The
CC sequences shown in ADD3224-ADD3260 represent murine mitochondrial DNA
CC clones used to make the probes of the invention. Some sequences are not
CC present, these are SEQ ID NO's 295, 1174, 1213, 1700, 1728, 1730, 1905,
CC 1906, 2408 and 2643.
XX
SQ Sequence 650 BP; 143 A; 172 C; 170 G; 165 T; 0 U; 0 Other;
XX

Alignment Scores:
Score: 3.21e-37 Length: 650
Pred. No.: 475.50 Matches: 94
Percent Similarity: 64.62% Conservative: 32
Best Local Similarity: 48.21% Mismatches: 56
Query Match: 43.87% Indels: 13
DB: 10 Gaps: 3

US-10-071-174-2 (1-204) x ADD34132 (1-650)
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QY 33 GluProGlyThrProGluProAlaProSerThrProGluAlaValLeuArgSerAla 52
Db 77 GAGCCGACACCCAGAGCCACCCGACCTGCGGAGCGGCTTGTCTCGCTCTGTG 136
QY 53 AlaAlaArgLeuArgGlnIleHisArgSerPhePheSerAlaTyrLeuGlyTyrProGly 72
Db 137 ACTAGGAGATCCAGAGAGGACCCACAGAAATTTTCTCTCTCGGAAAGCGGGGC 196
QY 73 AsnArgPheGluLeuValAlaLeuMetAlaAspSerValLeuSerAspSerProGlyPro 92
Db 197 AATCGCTGGAGCTGGTGAACACAGATGGCAGATAAGTTGCTCTCCAAAGACCAAGACTTC 256
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Db 257 AGCTGGAGCCAACTGGTGATGCTCTCTCGGCGGACGCTTATGAATCAAGGCGCT 316
QY 113 LeuValThrAlaArgTyrLysTyrGlyPheGlnProArgLeuLysGluGlnGly 132
Db 317 TACATGGCTGTCAAGCAGAAAGAGG-----GATCTGGG 349
QY 133 Asp-----ValAlaArgAspCysGlnArgLeuValAlaLeuLeuSerSerArgLeu 149
Db 350 AATCGTGTCTAGTACGCGGAGACTGCTCTCATAGTGAACCTTCTGTATAATCTGCTC 409
QY 150 MetGly---GlnHisArgAlaTrpLeuGlnAlaGlnGlyTrpAspGlyPheCysHis 168
Db 410 ATGGGCGTCCGACCCGCGCAGCTCGAGGCTCTCGCGCTGGGATGGCTTTTGGCGC 469
QY 169 PhePheArgThrProPheProLeuAlaPheTrpArgLysGlnLeuValGlnAlaPheLeu 188
Db 470 TTCTTCAAGATCTCTTACCGCTCTGGGAGAGATTGCTGATTCAAGGCTTTTCTG 529
QY 189 SerCysLeuLeuThrThrAlaPheIleTyrLeuTrpThrArgLeu 203
Db 530 TCAGGCTTCTTGAACAGCAACCTTTTATCTCGGAACGTTTA 574

RESULT 12
ADD34582/c
ID ADD34582 standard; DNA; 835 BP.
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AC ADD34582;
XX
DT 15-JAN-2004 (first entry)
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DE Mouse mitochondrial DNA sequence SEQ ID NO:2360.
XX
KW ds; mouse; array; mitochondrial; hybridisation; energy-metabolism;
KW mitochondrial disease; oxidative phosphorylation dysfunction;
KW oxidative stress; apoptosis; aging.
XX
OS Mus musculus.

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 7, 2004, 01:20:07 ; Search time 6234 Seconds
(without alignments)
6728.579 Million cell updates/sec

Title: US-10-071-174-1

Perfect score: 887

Sequence: 1 cgggccaagaaacacagcga.....ctctctcttgagtgaagaa 887

Scoring table: OLIGO NUC
Gapop_60.0 , Gapext 60.0

Searched: 4526729 seqs, 23644849745 residues

Word size : 0

Total number of hits satisfying chosen parameters: 9053458

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : GenEmbl.*

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- 2: gb_intg.*
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- 5: gb_ov.*
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- 7: gb_ph.*
- 8: gb_pl.*
- 9: gb_pr.*
- 10: gb_ro.*
- 11: gb_sts.*
- 12: gb_sy.*
- 13: gb_un.*
- 14: gb_vi.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	887	100.0	887	9 AF326964	AF326964 Homo sapi
2	763	86.0	1168	6 BD233466	BD233466 Human pro
3	615	69.3	615	9 AF285032	AF285032 Homo sapi
4	585	66.0	585	9 HS458330	AJ458330 Homo sapi
5	582	65.6	582	6 BD233456	BD233456 Human pro
C 6	539	60.8	93287	9 AC023906	AC023906 Homo sapi
C 7	439	49.5	726	6 CQ752105	CQ752105 Sequence
C 8	267	30.1	214669	2 AC018903	AC018903 Homo sapi
9	60	6.8	60	6 CQ545914	CQ545914 Sequence
10	47	5.3	214669	2 AC018903	AC018903 Homo sapi
C 11	22	2.5	5147	6 BD183336	BD183336 Novel gen
C 12	22	2.5	5147	6 BD171150	BD171150 Novel gen
C 13	22	2.5	5147	9 AB051441	AB051441 Homo sapi
14	22	2.5	21560	9 AY245248	AY245248 Homo sapi
15	22	2.5	60828	9 HS1191B2	AL022237 Human DNA
C 16	22	2.5	277861	2 HSAC000406	BC030863 Homo sapi
17	21	2.4	3011	10 BC030863	BC030863 Mus muscu
18	21	2.4	4353	10 AK173033	AK173033 Mus muscu
19	21	2.4	6190	10 BC043122	BC043122 Mus muscu

C 20	21	2.4	13635	1 AE005068	AE005068 Halobacte
C 21	21	2.4	95178	2 AL391556	AL391556 Homo sapi
C 22	21	2.4	120311	10 AC006945	AC006945 Mus muscu
23	21	2.4	155300	2 AL929459	AL929459 Danio rer
C 24	21	2.4	163574	2 AC138826	AC138826 Homo sapi
25	21	2.4	167273	10 AC083894	AC083894 Mus muscu
C 26	21	2.4	172144	2 AC136529	AC136529 Rattus no
C 27	21	2.4	172997	9 AC117516	AC117516 Homo sapi
C 28	21	2.4	190189	2 AC027181	AC027181 Homo sapi
C 29	21	2.4	224038	10 AL732625	AL732625 Mouse DNA
C 30	21	2.4	226457	2 AC127098	AC127098 Rattus no
C 31	21	2.4	231546	2 AC125302	AC125302 Rattus no
32	21	2.4	238290	2 AC112431	AC112431 Rattus no
33	21	2.4	288494	2 AC111749	AC111749 Rattus no
C 34	21	2.4	316892	2 AC096101	AC096101 Rattus no
C 35	20	2.3	284	11 G47512	G47512 Z25086_1 Ze
36	20	2.3	823	6 CQ780061	CQ780061 Sequence
37	20	2.3	823	6 CQ782081	CQ782081 Sequence
38	20	2.3	823	6 BD124770	BD124770 Primer fo
39	20	2.3	823	6 BD126790	BD126790 Primer fo
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41	20	2.3	1209	10 AF102501	AF102501 Mus muscu
42	20	2.3	1225	10 AF067660	AF067660 Mus muscu
43	20	2.3	1257	10 BC052690	BC052690 Mus muscu
C 44	20	2.3	1672	9 AY167728	AY167728 Pongo pyg
45	20	2.3	38674	9 HS081831	U81831 Human cosmi

ALIGNMENTS

RESULT 1 AF326964 887 bp mRNA linear PRI 01-MAY-2001
LOCUS Homo sapiens BCL2 (BCL2) mRNA, complete cds.
AF326964
DEFINITION AF326964.1 GI:13898393
VERSION AF326964.1
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 887)
AUTHORS Ke,N., Godzik,A. and Reed,J.C.
TITLE Bcl-2, a novel Bcl-2 family member that differentially binds and
regulates Bax and Bak
J. Biol. Chem. 276 (16), 12481-12484 (2001)
JOURNAL MEDLINE
PUBMED 21201065
REFERENCE 2 (bases 1 to 887)
AUTHORS Ke,N., Godzik,A. and Reed,J.C.
TITLE Direct Submission
SUBMITTED (07-DEC-2000) The Burnham Institute, 10901 N. Torrey
Pines Rd., La Jolla, CA 92037, USA
JOURNAL Location/Qualifiers
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Db	601	CTTTTAAACCGCTTCTACCTGCCCACTGFGACAACTAAATGACAGATGTGTGAGAAC	660
Qy	734	AGAACTGAGGAAAGACACCTTCCCAACCCACAGACGTTTTTATCTCAATGATACAAAGGA	793
Db	661	AGAACTGAGGAAAGACACCTTCCCAACCCACAGACGTTTTTATCTCAATGATACAAAGGA	720
Qy	794	GTCTGAGGTGGTGAATTTGCCAGTGTTTAACTGTGACAACTACTCAGGTGTGAGGAC	853
Db	721	GTCTGAGGTGGTGAATTTGCCAGTGTTTAACTGTGACAACTACTCAGGTGTGAGGAC	780
Qy	854	AGAAATCAAAATGGCTCTTCCCTTGGAGTGAAGAA	887
Db	781	AGAAATCAAAATGGCTCTTCCCTTGGAGTGAAGAA	814
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LOCUS	AF285092	615 bp mRNA linear	PRI 08-NOV-2001
DEFINITION	Homo sapiens Bcl-2-like protein 10 mRNA, complete cds.		
ACCESSION	AF285092		
VERSION	AF285092.1	GI:9837265	
KEYWORDS	Homo sapiens (human)		
SOURCE	Homo sapiens		
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
REFERENCE	1 (bases 1 to 615)		
AUTHORS	Zhang, H., Holmgren, W. and De Geyter, C.		
TITLE	Bcl2-L-10, a novel anti-apoptotic member of the Bcl-2 family, blocks apoptosis in the mitochondria death pathway but not in the death receptor pathway		
JOURNAL	Hum. Mol. Genet. 10 (21), 2329-2339 (2001)		
MEDLINE	21548034		
PUBMED	11689480		
REFERENCE	2 (bases 1 to 615)		
AUTHORS	Zhang, H.H.		
TITLE	Direct Submission		
JOURNAL	Submitted (05-JUL-2000) University Women's Hospital, Schanzenstra 46, Basel 4057, Switzerland		
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Db	61	CTGTTGCTGGCGGACTACTGGGTACTGCGCCCGGGAAACCGCGACCCCGAGCCGCG	120

Qy	170	CCATCCAGCCCGAGGCGCGCTGCTGGCTCCGCGGCCGAGTTACGCGAGATTAC	229
Db	121	CCATCCAGCCCGAGGCGCGCTGCTGGCTCCGCGGCCGAGTTACGCGAGATTAC	180
Qy	230	CGGTCTTTTCTCGGCTACCTCGGCTACCCCGGACCGCTTCGAGCTGGTGGCGCTG	289
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Qy	350	GTGACCTTCGAGGAGCGCTGCTGGAGAGAGGCGCTGTGTACCGCCCGGTGGAAGAAG	409
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Qy	410	TGGGCTTCCAGCCCGCTAAAGAGCAGGAGGCGGACGCTGCCCGGGACTGCCAGCGC	469
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Qy	470	CTGTTGGCTTCTGAGCTCGGCTCATGGGGCAGCAGCGCGCTGGCTGCGAGGCTCAG	529
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Qy	530	GGCGCTGGGATGCTTTTGTCACTTCTCAGGACCCCTTCCACTGGCTTTTGAGA	589
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Qy	590	AAACAGCTGGTCCAGGCTTTCTCTCATGTTGTATTAACAACAGCTTCATTATCTCTGG	649
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Qy	650	ACAGCATTTATTATGA	664
Db	601	ACAGCATTTATTATGA	615
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DEFINITION	Homo sapiens NRH gene for anti-apoptotic protein.		
ACCESSION	AJ458330		
VERSION	AJ458330.1	GI:20338765	
KEYWORDS	anti-apoptotic protein; NRH gene.		
SOURCE	Homo sapiens (human)		
ORGANISM	Homo sapiens		
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.		
REFERENCE	1		
AUTHORS	Aouacheria, A., Arnaud, E., Venet, S., Lalle, P., Gouy, M., Rigal, D. and Gillet, G.		
TITLE	Nrh, a human homologue of Nr-13 associates with Bcl-Xs and is an inhibitor of apoptosis		
JOURNAL	Oncogene 20 (41), 5846-5855 (2001)		
MEDLINE	21477277		
PUBMED	11593390		
REFERENCE	2 (bases 1 to 585)		
AUTHORS	Gillet, G.		
TITLE	Direct Submission		
JOURNAL	Submitted (23-APR-2002) Gillet G., Ibcop, CNRS UMR 5086, 7 PASSAGE DU VERCORS, 69367, FRANCE		
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Matches 585; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 140 GCCCGGGAACCCGACCCCGGAGCGGCCATCCACGCCGAGCGCGCGTGTGCGC 199
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DB 121 TCGCGGCGCCCGAGTTACGGCAGATTCACCGGTCTCTTTCTCGCCCTACCTCGGCTAC 180

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RESULT 5

BD233456
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DEFINITION BD233456
ACCESSION BD233456.1 GI:33043226
VERSION JP 2002519016-A/2
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

1 (bases 1 to 582)
Kato, S. and Kimura, T.
Human protein having hydrophobic domain and DNA encoding the same
Patent: JP 2002519016-A 2 02-JUL-2002;
JOURNAL SAGAMI CHEMICAL RESEARCH CENTER, PROTEGENE INC

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COMMENT OS Homo sapiens (human)
PN JP 2002519016-A/2
PD 02-JUL-2002
PF 18-JUN-1999 JP 2000557267
PI SEISHI KATO, TOMOKO KIMURA
PC C12N15/09, C07K14/47, C12N1/15, C12N1/19, C12N5/10, C12N15/00, C12N5/ PC
00 Human protein having hydrophobic domain and DNA encoding the
CC Human protein having hydrophobic domain and DNA encoding the
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DB 1 ATGGCCGACCCGCTGGGGAGCGCACCGAGCTGTGCTGGCCGACTACCTGGGGTACTGC 60

QY 140 GCCCGGGAACCCGACCCCGGAGCGGCCATCCACGCCGAGCGCGCGTGTGCGC 199
DB 61 GCCCGGGAACCCGACCCCGGAGCGGCCATCCACGCCGAGCGCGCGTGTGCGC 120

QY 200 TCGCGGCGCCCGAGTTACGGCAGATTCACCGGTCTCTTTCTCGCCCTACCTCGGCTAC 259
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DB 241 GGCCCGACCTGGGGCAGAGTGTGACGCTGTGACCTTCGAGGAGCGCTGTGGAGAGA 300

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DB 301 GGGCGCTGTGTGACCGCCGCTGCAAGTGTGGGCTTCAGCGCGGCTTAAAGGAGCAG 360

QY 440 GAGGCGACGCTGGCCGGGACTGCGAGCGCTGTGGCTTGTGAGCTTCGGGCTCATG 499
DB 361 GAGGCGACGCTGGCCGGGACTGCGAGCGCTGTGGCTTGTGAGCTTCGGGCTCATG 420

QY 500 GGGCAGACCGCGCTGGCTGCAAGTGTGGGCTTCAGCGCGGCTTAAAGGAGCAG 559
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QY 560 AGGACCCCTTTTCCACTGGCTTTTGGAGAAAACAGCTGTGTCAGGCTTTTGTGTCATGC 619
DB 481 AGGACCCCTTTTCCACTGGCTTTTGGAGAAAACAGCTGTGTCAGGCTTTTGTGTCATGC 540

QY 620 TTGTTAAACAACAGCGCTTCATTTATCTCTGGACAGATTATTA 661
DB 541 TTGTTAAACAACAGCGCTTCATTTATCTCTGGACAGATTATTA 582
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RESULT 6

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DEFINITION Homo sapiens chromosome 15 clone CTD-2184D3 map 15q21.2, complete
sequence.
ACCESSION AC023906
VERSION AC023906.7 GI:14595770
KEYWORDS HTG.

Center: Multimegabase Sequencing Center
Center code: UWMSC
Web site: <http://chroma.mbt.washington.edu/msg> www.


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ORIGIN
Query Match          6.8%; Score 60; DB 6; Length 60;
Best Local Similarity 100.0%; Pred. No. 3.2e-22;
Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 565 CCCTTTCACCTGGCTTTTGGAGAAAACAGCTGTGTCAGGCTTTTCGTGTCATGCTTGT 624
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Db 1 CCCTTTCACCTGGCTTTTGGAGAAAACAGCTGTGTCAGGCTTTTCGTGTCATGCTTGT 60

RESULT 10
AC018903      214669 bp DNA linear HTG 04-JUN-2000
LOCUS        Homo sapiens chromosome 15 clone RP11-337B11 map 15q21, LOW-PASS
DEFINITION   SEQUENCE SAMPLING.
ACCESSION   AC018903
VERSION      AC018903.2 GI:8247797
KEYWORDS     HTG; HTGS-PHASE0.
SOURCE       Homo sapiens (human)
ORGANISM     Homo sapiens

REFERENCE
AUTHORS      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
              1 (bases 1 to 214669)
              Rowen, L., Madan, A., Qin, S., Abbasi, N., Baradarani, L., Birditt, B.,
              Bloom, S., Dors, M., Dickhoff, R., Fleetwood, P., Harrison, G., Kaur, A.,
              Madan, A., Nesbitt, R., Shafer, T. and Hood, L.
              Sequencing of human chromosome 15 D15S146-D15S117 region
              Unpublished
              2 (bases 1 to 214669)
              Rowen, L., Madan, A., Qin, S., Abbasi, N., Baradarani, L., Birditt, B.,
              Bloom, S., Dors, M., Dickhoff, R., Fleetwood, P., Harrison, G.,
              James, R., Kaur, A., Madan, A., Owen, M. P., Ratcliffe, A., Shafer, T.
              and Hood, L.
              Direct Submission
              Submitted (22-DEC-1999) Multimegabase Sequencing Center, University
              of Washington, PO BOX 357730, Seattle, WA 98195, USA
              On Jun 4, 2000 this sequence version replaced gi:6630517.
              ----- Genome Center Sequencing Center
              Center: Multimegabase Sequencing Center
              Center code: UMSC
              Web site: http://chroma.mbt.washington.edu/msg_www
              Contact: leorowensystemsbiology.org
              ----- Summary Statistics
              Sequencing vector: pUC18; 108752
              Chemistry: Dye-terminator Big Dye; 90% of reads
              Chemistry: Dye-primer Big Dye; 10% of reads
              Assembly program: Phrap; version 0.990399
              -----
              * NOTE: This record contains 192 individual
              * sequencing reads that have not been assembled into
              * contigs. Runs of N are used to separate the reads
              * and the order in which they appear is completely
              * arbitrary. Low-pass sequence sampling is useful for
              * identifying clones that may be gene-rich and allows
              * overlap relationships among clones to be deduced.
              * However, it should not be assumed that this clone
              * will be sequenced to completion. In the event that
              * the record is updated, the accession number will
              * be preserved.
              *
              * 806: contig of 806 bp in length
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              * 907: contig of 1162 bp in length
              * 2068: gap of unknown length
              * 2168: contig of 852 bp in length
              * 3020: contig of 1164 bp in length
              * 3120: gap of unknown length
              * 3121: contig of 1164 bp in length
              * 3124: gap of unknown length
              * 4285: gap of unknown length
              * 4385: contig of 817 bp in length
              * 5201: gap of unknown length
              * 5302: contig of 1243 bp in length
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              * 6545: contig of 828 bp in length
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              * 7473: gap of unknown length
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              * 8857: contig of 802 bp in length
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              * 20800: contig of 1236 bp in length
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              * 25142: contig of 1144 bp in length
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  Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 512 GCCTGGCTGCAGGCTCAGGCG 533
Db 4330 GCCTGGCTGCAGGCTCAGGCG 4309

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DEFINITION Homo sapiens mRNA for KIAA1654 protein, partial cds.
ACCESSION AB051441
VERSION AB051441.1 GI:13359180
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Hirosewa,M., Nagase,T., Murhashi,Y., Kikuno,R. and Ohara,O.
TITLE Identification of novel transcribed sequences on human chromosome
JOURNAL DNA Res. 8 (1), 1-9 (2001)
MEDLINE 21156230
PUBMED 11258795
REFERENCE 2 (bases 1 to 5147)
AUTHORS Ohara,O., Nagase,T. and Kikuno,R.
TITLE Direct Submission
JOURNAL Submitted (22-NOV-2000) Osamu Ohara, Kazusa DNA Research Institute,
Department of Human Gene Research; 1532-3, Yana, Kisarazu, Chiba
292-0812, Japan (E-mail:cdna@kazusa.or.jp,
URL:http://www.kazusa.or.jp/huge, Tel:81-438-52-3913,
Fax:81-438-52-3914)

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  Best Local Similarity 100.0%; Pred. No. 2.5;
  Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 512 GCCTGGCTGCAGGCTCAGGCG 533
Db 4330 GCCTGGCTGCAGGCTCAGGCG 4309

RESULT 14

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AY245248
LOCUS Homo sapiens BCL2-interacting killer (apoptosis-inducing) (BIK)
DEFINITION Homo sapiens BCL2-interacting killer (apoptosis-inducing) (BIK)
ACCESSION AY245248
VERSION AY245248.1 GI:28932927
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 21560)
AUTHORS Rieder,M.J., Livingston,R.J., Daniels,M.R., Chung,M.-W.,
Miyamoto,K.E., Nguyen,C.P., Nguyen,D.A., Poel,C.L., Robertson,P.D.,
Schackwitz,W.S., Sherwood,J.K., Witrak,L.A. and Nickerson,D.A.
TITLE Direct Submission
JOURNAL Submitted (27-FEB-2003) Genome Sciences, University of Washington,
1705 NE Pacific, Seattle, WA 98195, USA
COMMENT To cite this work please use: NIEHS-SNPs, Environmental Genome
Project, NIEHS ES15478, Department of Genome Sciences, Seattle, WA
(URL: http://egp.gs.washington.edu).
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complement(8974. .9110)
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9733. .9746
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10640. .10765
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10853. .10899
/note="2.2 copies 21 mer GTGTGAGTACATGGGTGTCT 60%
conserved"
10887. .11068
/note="16.5 copies 11 mer GGTGTGTGTGT 114% conserved"
11135. .11150
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11215. .11521
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11523. .11886
/note="L1R10C repeat: matches 215. .562 of consensus"
11899. .11917
/note="9.5 copies 2 mer TG 29% conserved"
12600. .12893
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12894. .13201
/note="AluYc2 repeat: matches 1. .309 of consensus"
13205. .13220
/note="5.3 copies 3 mer AAG 32% conserved"
13224. .13241
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Best Local Similarity 100.0%; Pred. No. 2.8;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      512 GCCTGGCTCAGGCTCAGGCG 533
Db      33682 GCCTGGCTCAGGCTCAGGCG 33703
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Search completed: November 7, 2004, 04:47:20
Job time : 6238 secs

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GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: November 6, 2004, 23:36:01 ; Search time 686 Seconds

(without alignments)
6787.517 Million cell updates/sec

Title: US-10-071-174-1

Perfect score: 887

Sequence: 1 cgggccaagaaacacgacga.....ctcttccttgactgaagaa 887

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 4134886 seqs, 2624710521 residues

Word size : 0

Total number of hits satisfying chosen parameters: 8269772

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : N_Geneseq_23Sep04:*

- 1: Geneseqn1980s:*
- 2: Geneseqn1990s:*
- 3: Geneseqn2000s:*
- 4: Geneseqn2001as:*
- 5: Geneseqn2001bs:*
- 6: Geneseqn2002as:*
- 7: Geneseqn2002bs:*
- 8: Geneseqn2003as:*
- 9: Geneseqn2003bs:*
- 10: Geneseqn2003cs:*
- 11: Geneseqn2003ds:*
- 12: Geneseqn2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	887	100.0	887	6	AAD46683
2	763	86.0	1168	3	Aaz290049 Hydrophob
3	582	65.6	582	3	Aaz290039
4	488	55.0	522	4	Aah47022
5	225	25.4	548	4	ABK41913
6	225	25.4	548	9	ADBS5980
7	60	6.8	60	6	ABN42801
8	24	2.7	24	6	AAD46685
9	23	2.6	23	6	AAD46684
10	22	2.5	22	6	AAD46689
11	22	2.5	28	6	AAD46686
12	22	2.5	34	6	AAD46688
13	22	2.5	874	6	ABQ44403
14	22	2.5	874	6	ABQ44402
15	22	2.5	5147	6	ABN83959
16	21	2.4	30	6	AAD46687
17	21	2.4	3827	12	ADM66989
18	20	2.3	650	10	ADD34132
19	20	2.3	823	4	AAK31741
20	20	2.3	823	4	AAK93761
21	20	2.3	823	12	ADL28168

22	20	2.3	823	12	ADL30188
c	23	20	835	10	ADD34582
c	24	20	90798	12	ADP68859
25	19	2.1	65	3	ABN30228
26	19	2.1	387	6	ADF56850
c	27	19	507	11	ABD15534
28	19	2.1	665	6	ABT09478
29	19	2.1	665	10	ADG30917
30	19	2.1	665	12	ADG45503
c	31	19	936	11	ABD15622
32	19	2.1	1630	2	AAQ28270
33	19	2.1	1630	10	ABT41845
34	19	2.1	1779	5	AA578125
35	19	2.1	3419	4	AA911196
36	19	2.1	5561	6	AB192223
c	37	19	9588	10	ADG37082
c	38	19	28198	10	ADG37080
39	19	2.1	33053	6	ABQ67005
40	19	2.1	143391	10	ADL13648
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c	42	18	474	9	ACH18221
43	18	2.0	530	3	AAF15017
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ALIGNMENTS

RESULT 1

AAD46683
ID AAD46683 standard; DNA; 887 BP.

XX
AC AAD46683;

DT 27-JAN-2003 (first entry)

XX
DE Human Bcl-B DNA.

XX Human; Bcl-2; Bcl-B; therapy; apoptosis; cell degenerative disorder;
KW proliferative disorder; muscle degeneration; Alzheimer's disease; CJD;
KW Creutzfeldt-Jacob's disease; Machado-Joseph disease; MJD; transgenic;
KW Parkinson's disease; Huntington's disease; HD; spinocerebellar ataxia;
KW SCA; dentatorubropallidoluysian atrophy; DRPLA; Kennedy's disease;
KW stroke; ischaemia; head trauma; neoplasia; anticonvulsant; vulnery;
KW neurotropic; neuroprotective; cytostatic; immunosuppressive; vasotropic;
KW cerebroprotective; autoimmune disorder; chromosome 15; gene; ds.

XX Homo sapiens.

XX Key Location/Qualifiers
FH CDS 50..664
FT /*tag= a
/*product= "Human Bcl-B protein"

WO200272601-A2.

PD 19-SEP-2002.

XX 07-FEB-2002; 2002WO-US003547.

XX 07-FEB-2001; 2001US-0267156P.

XX 07-FEB-2002; 2002US-00071174.

(BURN-) BURNHAM INST.

Reed JC, Ke N, Godzik A;

WPI; 2002-723312/78.

DR P-PSDB; AAE29097.

XX New isolated or recombinant Bcl-B nucleic acids and polypeptides, for
PT treating a disorder associated with apoptosis, such as cell degenerative

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QY	140	GCCCGGAACCGGACCCCGAGCCGCGCCATCCAGCCGAGCCGCGCTGTGCGC 199
DB	61	GCCCGGAACCGGACCCCGAGCCGCGCCATCCAGCCGAGCCGCGCTGTGCGC 120
QY	200	TCGCGCGCCGCTTACCGGAGATTCACCGGTCCCTTTCTCCGCTACCTCGGCTAC 259
DB	121	TCGCGCGCCGCTTACCGGAGATTCACCGGTCCCTTTCTCCGCTACCTCGGCTAC 180
QY	260	CCCGGGAACCGCTTCGAGCTGGTGGCGCTGATGGCGGATTCGCTCTCCGACGCC 319
DB	181	CCCGGGAACCGCTTCGAGCTGGTGGCGCTGATGGCGGATTCGCTCTCCGACGCC 240
QY	320	GGCCCGACCTGGGACAGTGGTACGCTCTGTGACCTTCGACGAGCGCTGTGGAG 379
DB	241	GGCCCGACCTGGGACAGTGGTACGCTCTGTGACCTTCGACGAGCGCTGTGGAG 300
QY	380	GGGCGCGTGGTACCGCCGCTGGAAGAGTGGGGCTTCCAGCGCGGCTAAAGGAG 439
DB	301	GGGCGCGTGGTACCGCCGCTGGAAGAGTGGGGCTTCCAGCGCGGCTAAAGGAG 360
QY	440	GAGGGGACGCTGGCGGAGCTGCCAGCGCTGTGGCCCTGTGTGAGCTCGCGCTCAT 499
DB	361	GAGGGGACGCTGGCGGAGCTGCCAGCGCTGTGGCCCTGTGTGAGCTCGCGCTCAT 420
QY	500	GGGCGACACCGCGCTGGCTGACGCTCAGGCGGCTGGGATGGCTTTGTCACTTCT 559
DB	421	GGGCGACACCGCGCTGGCTGACGCTCAGGCGGCTGGGATGGCTTTGTCACTTCT 480
QY	560	AGGACCCCTTTCCACTGGCTTTTGGAGAAACAGCTGTCCAGGCTTTCTGTATGC 619
DB	481	AGGACCCCTTTCCACTGGCTTTTGGAGAAACAGCTGTCCAGGCTTTCTGTATGC 540
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DB	541	TTGTATACACAGCTTCATTATCTCTGGACAGGATTATTA 582
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XX	AAH47022 standard; cDNA; 522 BP.	
AC	AAH47022;	
DT	29-OCT-2001 (first entry)	
XX	Human Bcl-2-like polypeptide encoding cDNA (clone HLIBE40).	
DE	Bcl-2-like polypeptide; autoimmune disorder; allergy; immunomodulatory;	
KW	respiratory; cardiovascular; antiarthritic; immunostimulant; vaccine;	
KW	immunosuppressive; antiinflammatory; gene therapy; ss.	
XX	Homo sapiens.	
OS		
XX	Key	Location/Qualifiers
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FT		/note= "gene No. 2"
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PN	WO200157060-A1.	
XX		
PD	09-AUG-2001.	
XX		
PF	31-JAN-2001; 2001WO-US003080.	
XX		

PR	01-FEB-2000; 2000US-0179487P.	
PR	07-FEB-2000; 2000US-0180697P.	
XX	(HUMA-) HUMAN GENOME SCI INC.	
XX	Ruben SM, Duan DR, Ni J;	
PI	WPI; 2001-476279/51.	
XX	P-PSDB; AAB85666.	
DR	Nucleic acids encoding human Bcl-2-like polypeptides, useful for	
PT	preventing, diagnosing and/or treating.	
PT	Claim 1; Page 276; 285pp; English.	
PS	The invention provides nucleic acid molecules (NAM1) encoding 4 human Bcl	
XX	-2-like polypeptides (PEP1). The NAM1 and PEP1 may be used in the	
CC	prevention, diagnosis and treatment of diseases associated with	
CC	inappropriate Bcl-2-like polypeptides' expression. The NAM1 may be used	
CC	to produce the soluble Bcl-2-like polypeptides by standard recombinant	
CC	methodology. The polypeptides may also be used as antigens in the	
CC	production of antibodies against Bcl-2 and in assays to identify	
CC	modulators of Bcl-2 expression and activity. The anti-Bcl-2 antibodies	
CC	and antagonists may be used to down regulate expression and activity. The	
CC	anti-PEP1 antibodies may also be used as diagnostic agents for detecting	
CC	the presence of Bcl-2 polyps in samples (e.g. by enzyme linked	
CC	immunosorbant assay (ELISA)). Disorders that may be prevented, diagnosed	
CC	and/or treated by the above methods include, immunodeficiencies (e.g. a	
CC	gamma-globulinemia and B cell lymphoproliferative disorder), autoimmune	
CC	disorders (e.g. rheumatoid arthritis and Grave's disease), allergic	
CC	reactions, inflammations, respiratory diseases and cardiovascular	
CC	disorders (a full list of disorders is given in the specification). The	
CC	present sequence represents a human Bcl-2-like polypeptide encoding cDNA	
XX	Sequence 522 BP; 102 A; 148 C; 154 G; 118 T; 0 U; 0 Other;	
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QY	Query Match	55.0%; Score 488; DB 4; Length 522;
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QY	Matches 488; Conservative	0; Mismatches 0; Indels 0; Gaps 0;
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QY	330	GGGGCAGAGTGGTGAAGAGTGGGGCTTCCAGCGCGGCTAAAGGAGCAGAGGGCGGCG 389
DB	61	GGGGCAGAGTGGTGAAGAGTGGGGCTTCCAGCGCGGCTAAAGGAGCAGAGGGCGGCG 120
QY	390	TGACCGCCCGGTGGAAGAGTGGGGCTTCCAGCGCGGCTAAAGGAGCAGAGGGCGGCG 449
DB	121	TGACCGCCCGGTGGAAGAGTGGGGCTTCCAGCGCGGCTAAAGGAGCAGAGGGCGGCG 180
QY	450	TGCGCCGGGATGCGAGCGCTGTGGCTTGTGAGCTCGCGCTCATGGGGCAGACCC 509
DB	181	TGCGCCGGGATGCGAGCGCTGTGGCTTGTGAGCTCGCGCTCATGGGGCAGACCC 240
QY	510	GCGCTGCTGCTGAGCTCAGCGCGCTGGGATGGCTTTTGTCACTTCTTACGAGACCCCT 569
DB	241	GCGCTGCTGCTGAGCTCAGCGCGCTGGGATGGCTTTTGTCACTTCTTACGAGACCCCT 300
QY	570	TTCCACTGGCTTTTGGAGAAACAGCTGGTCCAGGCTTTTCTCATGCTGTGTAACA 629
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QY	630	CAGCTTTCATTATCTCTGGACACGATTAATATGAGTTTAAACTTTTAACTTCTTCT 689
DB	361	CAGCTTTCATTATCTCTGGACACGATTAATATGAGTTTAAACTTTTAACTTCTTCT 420
QY	690	ACCTGCCCAACTGTGACCAACTAAATGACAGATGTGTGAGAAACAAGAACTGAGGGAAGC 749
DB	421	ACCTGCCCAACTGTGACCAACTAAATGACAGATGTGTGAGAAACAAGAACTGAGGGAAGC 480
QY	750	ACCTTCCC 757

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Db      481 ACCTTCCC 488
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RESULT 5
ABK41913
ID   ABK41913 standard; cDNA; 548 BP.
XX
AC   ABK41913;
XX
XX 21-MAY-2002 (first entry)
XX
DE cDNA encoding novel human connective tissue related polypeptide #301.
XX
XX Human; connective tissue related disorder; cancer; gene therapy;
XX cytosstatic; gene; ss.
XX
XX Homo sapiens.
XX
XX WO200155343-A1.
XX
XX 02-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US001322.
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XX 31-JAN-2000; 2000US-0179065P.
XX 04-FEB-2000; 2000US-0180628P.
XX 24-FEB-2000; 2000US-0184664P.
XX 02-MAR-2000; 2000US-0186350P.
XX 16-MAR-2000; 2000US-0189874P.
XX 17-MAR-2000; 2000US-0190076P.
XX 18-APR-2000; 2000US-0198123P.
XX 19-MAY-2000; 2000US-0205515P.
XX 07-JUN-2000; 2000US-0209467P.
XX 28-JUN-2000; 2000US-0214886P.
XX 30-JUN-2000; 2000US-0215135P.
XX 07-JUL-2000; 2000US-0216647P.
XX 07-JUL-2000; 2000US-0216880P.
XX 11-JUL-2000; 2000US-0217487P.
XX 11-JUL-2000; 2000US-0217496P.
XX 14-JUL-2000; 2000US-0218290P.
XX 26-JUL-2000; 2000US-0220363P.
XX 26-JUL-2000; 2000US-0220364P.
XX 14-AUG-2000; 2000US-0224518P.
XX 14-AUG-2000; 2000US-0224519P.
XX 14-AUG-2000; 2000US-0225213P.
XX 14-AUG-2000; 2000US-0225214P.
XX 14-AUG-2000; 2000US-0225266P.
XX 14-AUG-2000; 2000US-0225267P.
XX 14-AUG-2000; 2000US-0225268P.
XX 14-AUG-2000; 2000US-0225270P.
XX 14-AUG-2000; 2000US-0225477P.
XX 14-AUG-2000; 2000US-0225757P.
XX 14-AUG-2000; 2000US-0225758P.
XX 18-AUG-2000; 2000US-0226279P.
XX 22-AUG-2000; 2000US-0226681P.
XX 22-AUG-2000; 2000US-0226868P.
XX 22-AUG-2000; 2000US-0227182P.
XX 23-AUG-2000; 2000US-0227009P.
XX 30-AUG-2000; 2000US-0228924P.
XX 01-SEP-2000; 2000US-0229287P.
XX 01-SEP-2000; 2000US-0229343P.
XX 01-SEP-2000; 2000US-0229344P.
XX 05-SEP-2000; 2000US-0229345P.
XX 05-SEP-2000; 2000US-0229509P.
XX 05-SEP-2000; 2000US-0229513P.
XX 06-SEP-2000; 2000US-0230437P.
XX 06-SEP-2000; 2000US-0230438P.
XX 08-SEP-2000; 2000US-0231242P.
XX 08-SEP-2000; 2000US-0231243P.
XX 08-SEP-2000; 2000US-0231244P.
XX 08-SEP-2000; 2000US-0231413P.
XX 08-SEP-2000; 2000US-0231414P.
XX 08-SEP-2000; 2000US-0232080P.
XX 08-SEP-2000; 2000US-0232081P.
XX 12-SEP-2000; 2000US-0231968P.
XX 14-SEP-2000; 2000US-0232397P.
XX 14-SEP-2000; 2000US-0232398P.
XX 14-SEP-2000; 2000US-0232399P.
XX 14-SEP-2000; 2000US-0232400P.
XX 14-SEP-2000; 2000US-0232401P.
XX 14-SEP-2000; 2000US-0233063P.
XX 14-SEP-2000; 2000US-0233064P.
XX 14-SEP-2000; 2000US-0233065P.
XX 21-SEP-2000; 2000US-0234223P.
XX 21-SEP-2000; 2000US-0234274P.
XX 25-SEP-2000; 2000US-0234997P.
XX 25-SEP-2000; 2000US-0234998P.
XX 26-SEP-2000; 2000US-0235484P.
XX 27-SEP-2000; 2000US-0235834P.
XX 27-SEP-2000; 2000US-0235836P.
XX 29-SEP-2000; 2000US-0236327P.
XX 29-SEP-2000; 2000US-0236367P.
XX 29-SEP-2000; 2000US-0236368P.
XX 29-SEP-2000; 2000US-0236369P.
XX 29-SEP-2000; 2000US-0236370P.
XX 02-OCT-2000; 2000US-0236802P.
XX 02-OCT-2000; 2000US-0237037P.
XX 02-OCT-2000; 2000US-0237038P.
XX 02-OCT-2000; 2000US-0237039P.
XX 13-OCT-2000; 2000US-0239935P.
XX 13-OCT-2000; 2000US-0239937P.
XX 20-OCT-2000; 2000US-0240960P.
XX 20-OCT-2000; 2000US-024121P.
XX 20-OCT-2000; 2000US-0241785P.
XX 20-OCT-2000; 2000US-0241786P.
XX 20-OCT-2000; 2000US-0241787P.
XX 20-OCT-2000; 2000US-0241808P.
XX 20-OCT-2000; 2000US-0241809P.
XX 01-NOV-2000; 2000US-0244617P.
XX 08-NOV-2000; 2000US-0246474P.
XX 08-NOV-2000; 2000US-0246475P.
XX 08-NOV-2000; 2000US-0246476P.
XX 08-NOV-2000; 2000US-0246477P.
XX 08-NOV-2000; 2000US-0246478P.
XX 08-NOV-2000; 2000US-0246523P.
XX 08-NOV-2000; 2000US-0246524P.
XX 08-NOV-2000; 2000US-0246525P.
XX 08-NOV-2000; 2000US-0246526P.
XX 08-NOV-2000; 2000US-0246527P.
XX 08-NOV-2000; 2000US-0246528P.
XX 08-NOV-2000; 2000US-0246532P.
XX 08-NOV-2000; 2000US-0246609P.
XX 08-NOV-2000; 2000US-0246610P.
XX 08-NOV-2000; 2000US-0246611P.
XX 08-NOV-2000; 2000US-0246613P.
XX 17-NOV-2000; 2000US-0249207P.
XX 17-NOV-2000; 2000US-0249208P.
XX 17-NOV-2000; 2000US-0249209P.
XX 17-NOV-2000; 2000US-0249210P.
XX 17-NOV-2000; 2000US-0249211P.
XX 17-NOV-2000; 2000US-0249212P.
XX 17-NOV-2000; 2000US-0249213P.
XX 17-NOV-2000; 2000US-0249214P.
XX 17-NOV-2000; 2000US-0249215P.
XX 17-NOV-2000; 2000US-0249216P.
XX 17-NOV-2000; 2000US-0249217P.
XX 17-NOV-2000; 2000US-0249218P.
XX 17-NOV-2000; 2000US-0249244P.
XX 17-NOV-2000; 2000US-0249245P.
XX 17-NOV-2000; 2000US-0249246P.
XX 17-NOV-2000; 2000US-0249265P.
XX 17-NOV-2000; 2000US-0249297P.
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PR 17-NOV-2000; 2000US-0249299P.
 PR 17-NOV-2000; 2000US-0249300P.
 PR 01-DEC-2000; 2000US-0250160P.
 PR 01-DEC-2000; 2000US-0250391P.
 PR 05-DEC-2000; 2000US-0251030P.
 PR 05-DEC-2000; 2000US-0251388P.
 PR 05-DEC-2000; 2000US-0256719P.
 PR 05-DEC-2000; 2000US-0251479P.
 PR 05-DEC-2000; 2000US-0251856P.
 PR 08-DEC-2000; 2000US-0251868P.
 PR 08-DEC-2000; 2000US-0251989P.
 PR 08-DEC-2000; 2000US-0251990P.
 PR 11-DEC-2000; 2000US-0254097P.
 PR 05-JAN-2001; 2001US-0259678P.
 XX (HUMA-) HUMAN GENOME SCI INC.
 PA
 XX Rosen CA, Barash SC, Ruben SM;
 XX
 XX WPI; 2001-565190/63.
 DR P-PSDB; AAU86735.
 XX
 XX Nucleic acid encoding novel connective tissue associated polypeptides,
 PT used in diagnosing, preventing, treating or ameliorating a disorder such
 PT as cancer or rheumatoid arthritis.
 PT
 XX Claim 4; SEQ ID NO 311; 673pp; English.
 PS
 XX The present invention relates to the isolation of novel human connective
 CC tissue related polypeptides (AAU86435-AAU86923) and the polynucleotide
 CC (cDNA and genomic) sequences encoding them. The sequences of the
 CC invention are useful in the diagnosis, treatment, prevention and/or
 CC prognosis of diseases associated with connective tissue(s), including
 CC cancer. The polynucleotide sequences of the invention are also useful in
 CC gene therapy. ABK41613-ABK42101 represent cDNA sequences encoding the
 CC novel human connective tissue related polypeptides. Note: The sequence
 CC data for this patent did not form part of the printed specification, but
 CC was obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 XX Sequence 548 BP; 99 A; 154 C; 167 G; 121 T; 0 U; 7 Other;
 SQ
 Query Match 25.4%; Score 225; DB 4; Length 548;
 Best Local Similarity 100.0%; Pred. No. 9.2e-101;
 Matches 225; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 524 GCTCAGGCGCGTGGATGGCTTTTGTCACTTCTCAGGACCCCTTCCACTGGCTTTT 583
 Db 324 GCTCAGGCGCGTGGATGGCTTTTGTCACTTCTCAGGACCCCTTCCACTGGCTTTT 383
 Qy 584 TGGAGAAAACAGCTGGTCCAGGCTTTTCTGTCATGCTTGTAAACACAGCTTCAATTAT 643
 Db 384 TGGAGAAAACAGCTGGTCCAGGCTTTTCTGTCATGCTTGTAAACACAGCTTCAATTAT 443
 Qy 644 CTCTGGACACGATTATTATGAGTTTAAACTTTTAAACCGCTTCTACCTGCCCACTGT 703
 Db 444 CTCTGGACACGATTATTATGAGTTTAAACTTTTAAACCGCTTCTACCTGCCCACTGT 503
 Qy 704 GACCACTAAATGACAGATGTGAGAACCAAGAACTGAGGGAAG 748
 Db 504 GACCACTAAATGACAGATGTGAGAACCAAGAACTGAGGGAAG 548
 RESULT 6
 ID ADB59580 standard; cDNA; 548 BP.
 XX
 AC ADB59580;
 XX
 XX 04-DEC-2003 (first entry)
 XX
 DE Connective tissue related polynucleotide #301.

XX cyrostatic; neuroprotective; nootropic; antiparkinsonian; cardiovascular;
 KW antiarteriosclerotic; immunosuppressive; antirheumatic; antiarthritic;
 KW antiinflammatory; antiallergic; antiasthmatic; dermatological;
 KW nephrotropic; virucide; fungicide; antibacterial; antiparasitic;
 KW gene therapy; ds; connective tissues disorder; rheumatoid arthritis;
 KW systemic lupus erythematosus; scleroderma; Sjogren's syndrome; cancer;
 KW cancer metastasis; neoplasia; leukemia; neurodegenerative disorder;
 KW Alzheimer's disease; Parkinson's disease; cardiovascular disease;
 KW atherosclerosis; myocarditis; cardiopulmonary bypass complication;
 KW autoimmune disease; multiple sclerosis; allergic reaction; asthma;
 KW rhinitis; eczema; inflammatory condition; Crohn's disease; nephritis;
 KW gastrointestinal disorder; inflammatory bowel disease;
 KW organ transplant rejection; immune system disorder; Bruton's disease;
 KW X-linked lymphoproliferative syndrome;
 KW B-cell lymphoproliferative disorder; HIV; AIDS; infection;
 KW chromosome identification; chromosome mapping;
 KW connective tissue related polynucleotide; gene; ss.
 XX Homo sapiens.
 XX
 XX US2003054375-A1.
 PN
 XX 20-MAR-2003.
 PD
 XX 07-MAR-2002; 2002US-00092154.
 PF
 XX 31-JAN-2000; 2000US-0179065P.
 PR 04-FEB-2000; 2000US-0180628P.
 PR 24-FEB-2000; 2000US-0184664P.
 PR 02-MAR-2000; 2000US-0186350P.
 PR 16-MAR-2000; 2000US-0189874P.
 PR 17-MAR-2000; 2000US-0190076P.
 PR 18-APR-2000; 2000US-0198123P.
 PR 19-MAY-2000; 2000US-0205151P.
 PR 27-JUN-2000; 2000US-0209467P.
 PR 28-JUN-2000; 2000US-0214866P.
 PR 30-JUN-2000; 2000US-0215135P.
 PR 07-JUL-2000; 2000US-0216647P.
 PR 07-JUL-2000; 2000US-0216880P.
 PR 11-JUL-2000; 2000US-0217487P.
 PR 11-JUL-2000; 2000US-0217496P.
 PR 14-JUL-2000; 2000US-0218290P.
 PR 26-JUL-2000; 2000US-0220963P.
 PR 26-JUL-2000; 2000US-0220964P.
 PR 14-AUG-2000; 2000US-0224518P.
 PR 14-AUG-2000; 2000US-0224519P.
 PR 14-AUG-2000; 2000US-0225213P.
 PR 14-AUG-2000; 2000US-0225214P.
 PR 14-AUG-2000; 2000US-0225266P.
 PR 14-AUG-2000; 2000US-0225287P.
 PR 14-AUG-2000; 2000US-0225288P.
 PR 14-AUG-2000; 2000US-0225270P.
 PR 14-AUG-2000; 2000US-0225447P.
 PR 14-AUG-2000; 2000US-0225757P.
 PR 14-AUG-2000; 2000US-0225758P.
 PR 14-AUG-2000; 2000US-0225759P.
 PR 18-AUG-2000; 2000US-0226279P.
 PR 22-AUG-2000; 2000US-0226681P.
 PR 22-AUG-2000; 2000US-0226865P.
 PR 22-AUG-2000; 2000US-0227182P.
 PR 23-AUG-2000; 2000US-0227009P.
 PR 30-AUG-2000; 2000US-0228924P.
 PR 01-SEP-2000; 2000US-0229287P.
 PR 01-SEP-2000; 2000US-0229343P.
 PR 01-SEP-2000; 2000US-0229344P.
 PR 01-SEP-2000; 2000US-0229345P.
 PR 05-SEP-2000; 2000US-0229509P.
 PR 05-SEP-2000; 2000US-0229513P.
 PR 06-SEP-2000; 2000US-0230437P.
 PR 06-SEP-2000; 2000US-0230438P.
 PR 08-SEP-2000; 2000US-0231242P.
 PR 08-SEP-2000; 2000US-0231243P.

AC ABN42801;
 XX
 DT 15-JUL-2002 (first entry)
 XX
 DE Human spliced transcript detection oligonucleotide SEQ ID NO:15549.
 XX
 KW Human; mouse; rat; splice transcript; detection; RNA transcript;
 KW splice variant; transcriptome; oligonucleotide library; ss.
 KW
 XX Homo sapiens.
 OS
 FN WO200210449-A2.
 XX
 PD 07-FEB-2002.
 XX
 PF 20-JUL-2001; 2001WO-1B001903.
 XX
 PR 28-JUL-2000; 2000US-0221607P.
 PR 02-MAY-2001; 2001US-0287724P.
 XX
 PA (COMP-) COMPUGEN INC.
 XX
 PI Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;
 XX
 DR WPI; 2002-257383/30.
 XX
 XX New oligonucleotide libraries comprising oligonucleotides which
 PT selectively hybridize to mRNAs transcribed from a transcription unit of a
 PT genome useful for detecting tissue-, pathology-, and developmental-
 PT specific genes.
 XX
 PS Example 1; SEQ ID NO 15549; 47pp; English.
 XX
 CC The present invention describes oligonucleotide libraries for detecting
 CC messenger RNAs that populate a (sub-)transcriptome, where the (sub-
 CC)transcriptome comprises messenger RNAs transcribed from multiple
 CC transcription units that populate a genome. The library comprises several
 CC oligonucleotides, each capable of hybridizing selectively to a set of
 CC messenger RNAs transcribed from a given transcription unit of the genome,
 CC which encodes one or more messenger RNA splice variants. The
 CC oligonucleotide libraries are useful for detecting mRNAs from a
 CC biological sample, in expression profiling studies, in qualitatively or
 CC quantitatively characterizing the corresponding transcriptome, and in
 CC detecting RNA transcripts and splice variants of human or animal
 CC transcriptomes. The libraries may also be used as specialized mini
 CC libraries to detect transcripts of a sub-transcriptome under a particular
 CC biological or pathological state, and so allowing the detection of tissue
 CC - and pathology-specific genes such as those genes only expressed in
 CC specific tissue under a specific pathological condition; to detect
 CC developmental specific genes; and to detect RNA transcripts and splice
 CC variants of a transcriptome of a patient suffering from a particular
 CC disorder. ABN27253 to ABN59589 represent oligonucleotide sequences from
 CC rats, humans and mice, which are used in the exemplification of the
 CC present invention. N.B. The sequence data for this patent did not form
 CC part of the printed specification, but was obtained in electronic format
 CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 60 BP; 9 A; 16 C; 13 G; 22 T; 0 U; 0 Other;
 XX
 Query Match 6.8%; Score 60; DB 6; Length 60;
 Best Local Similarity 100.0%; Pred. No. 4.4e-19;
 Matches 60; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 QY 565 CCCCTTCCACTGCTTTTGGAGAAAACAGCTGGTCCAGGCTTTTCTGTCATGCTTGT 624
 DB 1 CCCCTTCCACTGCTTTTGGAGAAAACAGCTGGTCCAGGCTTTTCTGTCATGCTTGT 60
 XX
 RESULT 8
 ID AAD46685/c
 XX AAD46685 standard; DNA; 24 BP.
 AC AAD46685;
 XX
 DT 27-JAN-2003 (first entry)
 XX
 DE Human Bcl-B cDNA cloning forward RT-PCR primer, NK01.
 XX
 KW Human; Bcl-2; Bcl-B; therapy; apoptosis; cell degenerative disorder;
 KW proliferative disorder; muscle degeneration; Alzheimer's disease; CJD;
 KW Creutzfeldt-Jacob's disease; Machado-Joseph disease; MJD; transgenic;
 KW Parkinson's disease; Huntington's disease; HD; spinocerebellar ataxia;
 KW SCA; dentatorubralpallidoluysian atrophy; DRPLA; Kennedy's disease;
 KW stroke; ischaemia; head trauma; neoplasia; anticonvulsant; vulnary;
 KW neurotropic; neuroprotective; cytostatic; immunosuppressive; vasotropic;
 KW cerebroprotective; autoimmune disorder; reverse transcription; RT; PCR;
 KW primer; ss.
 XX
 OS Homo sapiens.
 XX
 FN WO200272601-A2.
 XX
 PD 19-SEP-2002.
 XX
 PF 07-FEB-2002; 2002WO-US003547.
 XX
 PR 07-FEB-2001; 2001US-0267166P.
 PR 07-FEB-2002; 2002US-00071174.
 XX
 PA (BURN-) BURNHAM INST.
 XX
 PI Reed JC, Ke N, Godzik A;
 XX
 DR WPI; 2002-723312/78.
 XX
 XX New isolated or recombinant Bcl-B nucleic acids and polypeptides, for
 PT treating a disorder associated with apoptosis, such as cell degenerative
 PT or proliferative disorder e.g. cancer, Alzheimer's disease or Parkinson's
 PT disease.
 XX
 PS Example 1; Page 47; 82pp; English.
 XX
 CC The invention relates to human member of Bcl-2 family Bcl-B protein and
 CC its corresponding nucleic acid. Bcl-B is useful in treating a subject
 CC having or at risk of a disorder associated with apoptosis, such as cell
 CC degenerative or proliferative disorder like neural or muscle
 CC degeneration, e.g. Alzheimer's disease, Creutzfeldt-Jacob's disease
 CC (CJD), Machado-Joseph disease (MJD), Parkinson's disease, Huntington's
 CC disease (HD), spinocerebellar ataxias 1, 2 and 6 (SCA-1, -2 and -6),
 CC dentatorubralpallidoluysian atrophy (DRPLA), Kennedy's disease, stroke,
 CC ischaemia, head trauma, neoplasia, autoimmune disorder or fibrotic
 CC condition. The transgenic animals are used as in vivo models to study
 CC apoptosis and potential therapies for apoptosis. The present sequence is
 CC a reverse transcription (RT)-PCR primer used for cloning human Bcl-B cDNA
 XX
 SQ Sequence 24 BP; 8 A; 6 C; 5 G; 5 T; 0 U; 0 Other;
 XX
 Query Match 2.7%; Score 24; DB 6; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.3;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 QY 858 ATGCAATGGCTCTTCCTTGAGTG 881
 DB 24 ATGCAATGGCTCTTCCTTGAGTG 1
 XX
 RESULT 9
 ID AAD46684
 XX AAD46684 standard; DNA; 23 BP.
 AC AAD46684;
 XX
 DT 27-JAN-2003 (first entry)
 XX
 DE Human Bcl-B cDNA cloning forward RT-PCR primer, NK01.
 XX
 KW Human Bcl-B cDNA cloning forward RT-PCR primer, NK01.

XX
 DT 27-JAN-2003 (first entry)
 XX
 DE Human Bcl-B cDNA cloning reverse RT-PCR primer, NK0121.
 XX
 KW Human; Bcl-2; Bcl-B; therapy; apoptosis; cell degenerative disorder;
 KW proliferative disorder; muscle degeneration; Alzheimer's disease; CJD;
 KW Creutzfeldt-Jacob's disease; Machado-Joseph disease; MJD; transgenic;
 KW Parkinson's disease; Huntington's disease; HD; spinocerebellar ataxia;
 KW SCA; dentatorubralpallidoluysian atrophy; DRPLA; Kennedy's disease;
 KW stroke; ischaemia; head trauma; neoplasia; anticonvulsant; vulnary;
 KW neurotropic; neuroprotective; cytostatic; immunosuppressive; vasotropic;
 KW cerebroprotective; autoimmune disorder; reverse transcription; RT; PCR;
 KW primer; ss.
 XX
 OS Homo sapiens.
 XX
 FN WO200272601-A2.
 XX
 PD 19-SEP-2002.
 XX
 PF 07-FEB-2002; 2002WO-US003547.
 XX
 PR 07-FEB-2001; 2001US-0267166P.
 PR 07-FEB-2002; 2002US-00071174.
 XX
 PA (BURN-) BURNHAM INST.
 XX
 PI Reed JC, Ke N, Godzik A;
 XX
 DR WPI; 2002-723312/78.
 XX
 XX New isolated or recombinant Bcl-B nucleic acids and polypeptides, for
 PT treating a disorder associated with apoptosis, such as cell degenerative
 PT or proliferative disorder e.g. cancer, Alzheimer's disease or Parkinson's
 PT disease.
 XX
 PS Example 1; Page 47; 82pp; English.
 XX
 CC The invention relates to human member of Bcl-2 family Bcl-B protein and
 CC its corresponding nucleic acid. Bcl-B is useful in treating a subject
 CC having or at risk of a disorder associated with apoptosis, such as cell
 CC degenerative or proliferative disorder like neural or muscle
 CC degeneration, e.g. Alzheimer's disease, Creutzfeldt-Jacob's disease
 CC (CJD), Machado-Joseph disease (MJD), Parkinson's disease, Huntington's
 CC disease (HD), spinocerebellar ataxias 1, 2 and 6 (SCA-1, -2 and -6),
 CC dentatorubralpallidoluysian atrophy (DRPLA), Kennedy's disease, stroke,
 CC ischaemia, head trauma, neoplasia, autoimmune disorder or fibrotic
 CC condition. The transgenic animals are used as in vivo models to study
 CC apoptosis and potential therapies for apoptosis. The present sequence is
 CC a reverse transcription (RT)-PCR primer used for cloning human Bcl-B cDNA
 XX
 SQ Sequence 24 BP; 8 A; 6 C; 5 G; 5 T; 0 U; 0 Other;
 XX
 Query Match 2.7%; Score 24; DB 6; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.3;
 Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 XX
 QY 858 ATGCAATGGCTCTTCCTTGAGTG 881
 DB 24 ATGCAATGGCTCTTCCTTGAGTG 1
 XX
 RESULT 9
 ID AAD46684
 XX AAD46684 standard; DNA; 23 BP.
 AC AAD46684;
 XX
 DT 27-JAN-2003 (first entry)
 XX
 DE Human Bcl-B cDNA cloning forward RT-PCR primer, NK01.

KW Human; Bcl-2; Bcl-B; therapy; apoptosis; cell degenerative disorder;
 KW proliferative disorder; muscle degeneration; Alzheimer's disease; CJD;
 KW Creutzfeldt-Jacob's disease; Machado-Joseph disease; MJD; transgenic;
 KW Parkinson's disease; Huntington's disease; HD; spinocerebellar ataxia;
 KW SCA; dentatorubropallidoluysian atrophy; DRPLA; Kennedy's disease;
 KW stroke; ischaemia; head trauma; neoplasia; anticonvulsant; vulnary;
 KW neurotropic; neuroprotective; cytosolic; immunosuppressive; vasotropic;
 KW cerebroprotective; autoimmune disorder; reverse transcription; RT; PCR;
 KW primer; ss.
 OS Homo sapiens.
 XX WO200272601-A2.
 PN 19-SEP-2002.
 XX 07-FEB-2002; 2002WO-US003547.
 XX 07-FEB-2001; 2001US-0267166P.
 PR 07-FEB-2002; 2002US-00071174.
 XX (BURN-) BURNHAM INST.
 PA Reed JC, Ke N, Godzik A;
 PI WPI; 2002-723312/78.
 DR New isolated or recombinant Bcl-B nucleic acids and polypeptides, for
 PT treating a disorder associated with apoptosis, such as cell degenerative
 PT or proliferative disorder e.g. cancer, Alzheimer's disease or Parkinson's
 PT disease.
 XX Example 1; Page 47; 82pp; English.
 PS The invention relates to human member of Bcl-2 family Bcl-B protein and
 CC its corresponding nucleic acid. Bcl-B is useful in treating a subject
 CC having or at risk of a disorder associated with apoptosis, such as cell
 CC degenerative or proliferative disorder like neural or muscle
 CC degeneration, e.g. Alzheimer's disease, Creutzfeldt-Jacob's disease
 CC (CJD), Machado-Joseph disease (MJD), Parkinson's disease, Huntington's
 CC disease (HD), spinocerebellar ataxia 1, 2 and 6 (SCA-1, -2 and -6),
 CC dentatorubropallidoluysian atrophy (DRPLA), Kennedy's disease, stroke,
 CC ischaemia, head trauma, neoplasia, autoimmune disorder or fibrotic
 CC condition. The transgenic animals are used as in vivo models to study
 CC apoptosis and potential therapies for apoptosis. The present sequence is
 CC a reverse transcription (RT)-PCR primer used for cloning human Bcl-B cDNA
 CC SQA Sequence 23 BP; 9 A; 6 C; 8 G; 0 T; 0 U; 0 Other;
 SQ Query Match 2.8%; Score 23; DB 6; Length 23;
 Best Local Similarity 100.0%; Pred. No. 0.94;
 Matches 23; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 CGGGCCCAAGAAACACCGAAGG 23
 DB 1 CGGGCCCAAGAAACACCGAAGG 23
 RESULT 10
 AAD46689
 ID AAD46689 standard; DNA; 22 BP.
 XX AAD46689;
 AC AAD46689;
 XX 27-JAN-2003 (first entry)
 DT Human Bcl-B cDNA amplifying forward RT-PCR primer, NK0120.
 DE Human; Bcl-2; Bcl-B; therapy; apoptosis; cell degenerative disorder;
 KW proliferative disorder; muscle degeneration; Alzheimer's disease; CJD;
 KW Creutzfeldt-Jacob's disease; Machado-Joseph disease; MJD; transgenic;
 KW Parkinson's disease; Huntington's disease; HD; spinocerebellar ataxia;
 KW SCA; dentatorubropallidoluysian atrophy; DRPLA; Kennedy's disease;
 KW stroke; ischaemia; head trauma; neoplasia; anticonvulsant; vulnary;
 KW neurotropic; neuroprotective; cytosolic; immunosuppressive; vasotropic;
 KW cerebroprotective; autoimmune disorder; reverse transcription; RT; PCR;
 KW primer; ss.

KW stroke; ischaemia; head trauma; neoplasia; anticonvulsant; vulnary;
 KW neurotropic; neuroprotective; cytosolic; immunosuppressive; vasotropic;
 KW cerebroprotective; autoimmune disorder; reverse transcription; RT; PCR;
 KW primer; ss.
 OS Homo sapiens.
 XX WO200272601-A2.
 PN 19-SEP-2002.
 XX 07-FEB-2002; 2002WO-US003547.
 XX 07-FEB-2001; 2001US-0267166P.
 PR 07-FEB-2002; 2002US-00071174.
 XX (BURN-) BURNHAM INST.
 PA Reed JC, Ke N, Godzik A;
 PI WPI; 2002-723312/78.
 DR New isolated or recombinant Bcl-B nucleic acids and polypeptides, for
 PT treating a disorder associated with apoptosis, such as cell degenerative
 PT or proliferative disorder e.g. cancer, Alzheimer's disease or Parkinson's
 PT disease.
 XX Example 1; Page 48; 82pp; English.
 PS The invention relates to human member of Bcl-2 family Bcl-B protein and
 CC its corresponding nucleic acid. Bcl-B is useful in treating a subject
 CC having or at risk of a disorder associated with apoptosis, such as cell
 CC degenerative or proliferative disorder like neural or muscle
 CC degeneration, e.g. Alzheimer's disease, Creutzfeldt-Jacob's disease
 CC (CJD), Machado-Joseph disease (MJD), Parkinson's disease, Huntington's
 CC disease (HD), spinocerebellar ataxia 1, 2 and 6 (SCA-1, -2 and -6),
 CC dentatorubropallidoluysian atrophy (DRPLA), Kennedy's disease, stroke,
 CC ischaemia, head trauma, neoplasia, autoimmune disorder or fibrotic
 CC condition. The transgenic animals are used as in vivo models to study
 CC apoptosis and potential therapies for apoptosis. The present sequence is
 CC a reverse transcription (RT)-PCR primer used for amplifying human Bcl-B
 CC cDNA
 CC SQA Sequence 22 BP; 2 A; 6 C; 8 G; 6 T; 0 U; 0 Other;
 SQ Query Match 2.5%; Score 22; DB 6; Length 22;
 Best Local Similarity 100.0%; Pred. No. 2.9;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 338 GTGGTGACGCTCGTGACCTTCG 359
 DB 1 GTGGTGACGCTCGTGACCTTCG 22
 RESULT 11
 AAD46686
 ID AAD46686 standard; DNA; 28 BP.
 XX AAD46686;
 AC AAD46686;
 XX 27-JAN-2003 (first entry)
 DT Human Bcl-B DNA amplifying forward PCR primer, NK0101.
 DE Human; Bcl-2; Bcl-B; therapy; apoptosis; cell degenerative disorder;
 KW proliferative disorder; muscle degeneration; Alzheimer's disease; CJD;
 KW Creutzfeldt-Jacob's disease; Machado-Joseph disease; MJD; transgenic;
 KW Parkinson's disease; Huntington's disease; HD; spinocerebellar ataxia;
 KW SCA; dentatorubropallidoluysian atrophy; DRPLA; Kennedy's disease;
 KW stroke; ischaemia; head trauma; neoplasia; anticonvulsant; vulnary;
 KW neurotropic; neuroprotective; cytosolic; immunosuppressive; vasotropic;
 KW cerebroprotective; autoimmune disorder; PCR; primer; ss.

XX Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons
PT from chemically treated DNA.
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridisation to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridised to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABQ13410-
CC ABQ54121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention
XX
SQ Sequence 874 BP; 374 A; 291 C; 90 G; 119 T; 0 U; 0 Other;
XX
Query Match 2.5%; Score 22; DB 6; Length 874;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 233 TCCTTTTCTCCGCTACCTCG 254
DB 332 TCCTTTTCTCCGCTACCTCG 353
XX
RESULT 14
ABQ4402/c
ID ABQ4402 standard; DNA; 874 BP.
AC ABQ44402;
XX
XX 12-JUL-2002 (first entry)
DE
XX Oligonucleotide for detecting cytosine methylation SEQ ID NO 30993.
XX Human; cytosine methylation; 5'-CpG-3'; uracil; cytosine; diagnosis;
KW drug; side effect; cancer; central nervous system; cardiovascular;
KW gastrointestinal; respiratory system; single nucleotide polymorphism;
KW SNP; cell differentiation; ds.
XX
XX Homo sapiens.
XX
XX WO200218632-A2.
XX
XX 07-MAR-2002.
XX
XX 01-SEP-2001; 2001WO-EP010074.
XX
XX 01-SEP-2000; 2000DE-01043826.
XX
XX 05-SEP-2000; 2000DE-01044543.
XX
XX (EPIG-) EPIGENOMICS AG.
XX
XX Olek A, Piepenbrock C, Berlin K, Guetig D;
XX WPI; 2002-371829/40.
XX
XX Determining the degree of cytosine methylation in genomic DNA, useful for
PT diagnosis and prognosis, comprises selective hybridization of amplicons

PT from chemically treated DNA.
XX
PS Claim 12; 56pp + Sequence Listing; 56pp; German.
XX
XX This invention describes a novel method for determining the degree of
CC methylation of a particular cytosine in a motif 5'-CpG-3', present in a
CC genomic sample of DNA. The sample is treated chemically to convert
CC cytosine (C) but not methylated C, to uracil, then part of the genomic
CC DNA that contains the target C is amplified to form a labeled amplicon.
CC The amplicon is hybridised to two classes, each with at least one member,
CC of oligonucleotides and/or peptide-nucleic acid (PNA) oligomers and the
CC degree of hybridisation to both classes is determined from the label on
CC the amplicon. From the ratio of labels hybridised to the two classes of
CC oligomers, the degree of methylation is calculated. The method is used:
CC (i) for diagnosis and/or prognosis of side effects of therapeutic drugs
CC and of a wide range of diseases, e.g. cancer, disorders of the central
CC nervous, cardiovascular, gastrointestinal and respiratory systems etc.,
CC particularly by detecting mutations or single nucleotide polymorphisms
CC (SNP's); and (ii) for differentiation of cell or tissue types and for
CC investigating cell differentiation. The method allows the methylation
CC status of many C residues to be determined simultaneously. ABQ13410-
CC ABQ54121 represent genomic DNA sequences used to illustrate the method
CC for determining the degree of cytosine methylation described in the
CC disclosure of the invention
XX
SQ Sequence 874 BP; 119 A; 90 C; 291 G; 374 T; 0 U; 0 Other;
XX
Query Match 2.5%; Score 22; DB 6; Length 874;
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 233 TCCTTTTCTCCGCTACCTCG 254
DB 543 TCCTTTTCTCCGCTACCTCG 522
XX
RESULT 15
ABN83959/c
ID ABN83959 standard; DNA; 5147 BP.
XX
AC ABN83959;
XX
XX 06-SEP-2002 (first entry)
DE
XX Human gene sequence #6.
XX Human; brain; tonsil; hippocampus; foetal brain; diagnosis; gene; ds.
XX
XX Homo sapiens.
XX
XX Key Location/Qualifiers
FT CDS 2074..2349
FT /*tag=a
FT /partial
FT /note= "no start codon present"
XX
XX WO200252005-A1.
XX
XX 04-JUL-2002.
XX
XX 20-DEC-2001; 2001WO-JP011217.
XX
XX 22-DEC-2000; 2000JP-00389742.
XX
XX (KAZU-) KAZUSA DNA RES INST FOUND.
XX (CELE-) CELESTAR LEXICO-SCI LTD.
XX
XX Ohara O, Nagase T, Nakajima D;
XX WPI; 2002-500762/53.
XX P-PSDB; ABB97939.
XX
XX Genes and their expression products cloned from human cDNA libraries for

PT treatment and diagnosis of diseases associated with their expression.
 XX
 PS Claim 1(a); Page 63-66; 238pp; Japanese.
 XX
 CC The invention relates to DNA encoding polypeptides directly cloned from
 CC cDNA libraries originating in adult whole brain, human tonsil, human
 CC adult hippocampus and human foetal brain. Polypeptides and
 CC polynucleotides of the invention may be used in the investigation of
 CC differential expression of the DNA sequences in normal subjects and
 CC disease patients. They may also be used in the production of antibodies,
 CC oligonucleotide probes and DNA chips for diagnosis and identification of
 CC drugs for treatment of diseases with which the DNA sequences are
 CC associated. The sequences given in records ABN83954-ABN83984 represent
 CC human gene sequences of the invention
 XX
 SQ Sequence 5147 BP; 1212 A; 1455 C; 1334 G; 1146 T; 0 U; 0 Other;
 Query Match 2.5%; Score 22; DB 6; Length 5147;
 Best Local Similarity 100.0%; Pred. No. 2.4;
 Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 512 GCCTGGCTGCAGGCTCAGGCG 533
 Db 4330 GCCTGGCTGCAGGCTCAGGCG 4309
 Search completed: November 7, 2004, 03:03:12
 Job time : 688 secs

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: November 7, 2004, 02:37:47 ; Search time 4548 Seconds
(without alignments)
7106.869 Million cell updates

Title: US-10-071-174-1

Perfect score: 887
Sequence: 1 cgggccaagaaaaaccagcga.....ctctccttgagtgaagaaa 887

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 32822875 seqs, 18219865908 residues

Word size : 0

Total number of hits satisfying chosen parameters: 65645750

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : EST:★

1: d_ess1: *

2: d_ess2: *

3: d_ess3: *

4: d_ess4: *

5: d_ess5: *

6: d_ess6: *

7: d_ess7: *

8: d_ess8: *

9: d_ess9: *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query			DB	ID	Description
	Match	Length	%			
C 1	197	22.2	535	6	CA777633	CA777633 ip19a09.y
C 2	167	18.8	515	5	EX119239	EX119239 BX119239.y
C 3	111	12.5	493	1	A1813346	A1813346 wj35903.x
C 4	91	10.3	130	1	AA098865	AA098865 zk84f02.s
C 5	53	6.0	206	1	AA005293	AA005293 zh93a11.r
C 6	30	3.4	478	7	R53538	R53538 yG94c01.s1
C 7	27	3.0	939	9	CNS02C0P	AL190546 Tetraodon
C 8	22	2.5	316	7	R85550	R85550 yo38d07.s1
C 9	22	2.5	601	8	BZ869242	BZ869242 CH240_211
C 10	21	2.4	441	8	AZ617122	AZ617122 LM0448D17
C 11	21	2.4	462	5	BY590905	BY590905 BY590905
C 12	21	2.4	479	8	AQ294438	AQ294438 HS_3014.B
C 13	21	2.4	494	1	AA823641	AA823641 vF59a10_.s
C 14	21	2.4	657	2	B8628603	B8628603 BB628603
C 15	21	2.4	803	9	CNS0413S	AL269713 Tetraodon
C 16	21	2.4	881	2	BF532088	BF532088 602073207
C 17	21	2.4	931	4	BG173025	BG173025 602335506
C 18	21	2.4	2780	3	AA035932	AA035932 Mus muscu
C 19	20	2.3	352	1	AK426934	AK426934 vF22b11_.s
C 20	20	2.3	415	5	BY059827	BY059827 BY059827
C 21	20	2.3	448	6	CA560084	CA560084 K0267E03-
C 22	20	2.3	454	1	AJ631643	AJ631643 AJ631643
C 23	20	2.3	455	8	AQ457939	AQ457939 HS_5189.B
C 24	20	2.3	478	6	CA561864	CA561864 K0293B04-

25	20	2.3	524	6	CA560564
26	20	2.3	545	4	BG071824
27	20	2.3	586	6	BG073550
28	20	2.3	596	7	CO702056
29	20	2.3	613	5	BP464181
30	20	2.3	626	4	BG080862
31	20	2.3	649	7	CF915355
32	20	2.3	710	8	BZ006250
33	20	2.3	728	9	EX969107
34	20	2.3	736	6	EY735639
35	20	2.3	758	7	CO799635
36	20	2.3	763	7	CO808612
37	20	2.3	772	7	CO814648
38	20	2.3	783	8	BZ452477
39	20	2.3	791	5	BU941870
40	20	2.3	800	9	CL351188
41	20	2.3	821	7	CO806531
42	20	2.3	852	8	BZ557066
43	20	2.3	875	7	CO798715
44	20	2.3	893	7	CO811958
45	19	2.1	190	4	BI301953

ALIGNMENTS

RESULT 1	CA777633/c	715 bp	mrna	linear	EST 03-DEC-2002
LOCUS	CA777633				
DEFINITION	ip19a09.y1 HR85 islet Homo sapiens cDNA clone IMAGE16217625 5' similar to TR:Q9Z0F3 Q9Z0F3 BCL-2 HOMOLOG., mRNA sequence.				
ACCESSION	CA777633				
VERSION	CA777633.1	GI:26015508			
KEYWORDS	EST.				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
REFERENCE	1. (bases 1 to 715)				
AUTHORS	Melton,D., Brown,J., Kenty,G., Permutt,A., Lee,C., Kaestner,K., Lemishka,I., Secearc,M., Brestelli,J., Gradwohl,G., Clifton,S., Hillier,L., Marra,M., Pape,D., Wylie,T., Martin,J., Blistain,A., Schmitt,A., Theising,B., Ritter,E., Ronko,I., Bennett,J., Cardenas,M., Gibbons,M., McCann,R., Cole,R., Tsagareishvili,R., Williams,T., Jackson,Y. and Bowers,Y.				
TITLE	Endocrine Pancreas Consortium				
JOURNAL	Unpublished (2000)				
COMMENT	Contact: Douglas Melton, Klaus H. Kaestner, & Hiroshi Inoue				

Harvard University, Howard Hughes Medical Institute
Dept of Molecular and Cellular Biology, 7 Divinity Ave, Cambridge,
MA 02138
Tel: 617-495-1812
Fax: 617-495-8557
Email: dmelton@biochp.harvard.edu
Library was constructed by Dr. Hiroshi Inoue DNA sequencing by:
Washington University Genome Sequencing Center For information on
obtaining a clone please contact: Dr. Hiroshi Inoue
(hinoue@im.wustl.edu)
Possible reversed clone: similarity on wrong strand
Seq primer: -40RP from Gibco
High quality sequence stop:392.

FEATURES
source

```
1. /r11
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/cdones="IMAGE:6217625"
/tissue_type="Purified pancreatic islet"
/lab_host="D110B"
/clone_lib="pHR8 islet"
/note="Organ: Pancreas; Vector: pBluecrd
Not1; Site 2: XhoI; cDNA made by oligo-d
```

Size-selected on agarose gel. Average insert size ~1kb. 5' XhoI site was destroyed after directional cloning. Amplified once. Contact information: Hiroshi Inoue, MD, Metabolism Div. (Alan Permutt Lab), Washington University School of Medicine, Box 8127, 660 South Euclid Ave., St. Louis, MO 63110, E-mail: hinoue@imgate.wustl.edu, Tel: 314-362-1916, Fax: 314-747-2692."

ORIGIN

Query Match 22.2%; Score 197; DB 6; Length 715;
 Best Local Similarity 100.0%; Pred. No. 5.6e-95;
 Matches 197; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 691 CTTGCCCACTGTGACCACTAAATGACAGATGTGTGAGAACAACTGAGGAAAGCA 750
 Db 581 CTTGCCCACTGTGACCACTAAATGACAGATGTGTGAGAACAACTGAGGAAAGCA 522
 QY 751 CTTTCCCCCACCAGACGTTTTATCTGAATGCATACAGGAGTCTCGAGGTGGTATT 810
 Db 521 CTTTCCCCCACCAGACGTTTTATCTGAATGCATACAGGAGTCTCGAGGTGGTATT 462
 QY 811 TGGCCAGTGTTTAACTTTGTGACAACTGACAGTGTGAGGACAAAGTGCAAATGGCTC 870
 Db 461 TGGCCAGTGTTTAACTTTGTGACAACTGACAGTGTGAGGACAAAGTGCAAATGGCTC 402
 QY 871 TTTCTTCACTGAAGAA 887
 Db 401 TTTCTTCACTGAAGAA 385

RESULT 2

BX119239/c
 LOCUS BX119239 535 bp mRNA linear EST 10-FEB-2003
 DEFINITION BX119239 Soares infant brain IN1B Homo sapiens cDNA clone
 IMAGP998N01170 ; IMAGE:40052, mRNA sequence.
 ACCESSION BX119239
 VERSION BX119239.1 GI:27841652
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 AUTHORS Ebert L., Heil O., Hennig S., Neubert P., Partsch E., Peters M.,
 Radelof U., Schneider D. and Korn B.
 TITLE Human Unigeneset - RZPD3
 JOURNAL Unpublished (2003)
 COMMENT Contact: Ina Rolfs

RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
 Im Neuenheimer Feld 580, D-69120 Heidelberg, Germany
 RZPDLB; I.M.A.G.E. cDNA Clone Collection;
 Human Unigeneset - RZPD3 (RZPDLB No.972)
 http://www.rzpd.de/CloneCards/cgi-bin/showlib.pl.cgi/response?libNo=972 Contact: Ina Rolfs
 RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
 Heubnerweg 6, D-14059 Berlin, Germany
 Tel: +49 30 32639 101
 Fax: +49 30 32639 111
 www.rzpd.de

This clone is available royalty-free from RZPD;
 contact RZPD (clone@rzpd.de) for further information. Seq primer:
 M13u, Primer sequence: CGTGTAAACACGCGCCAGT.

FEATURES

source
 1..535
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGP998N01170 ; IMAGE:40052"
 /sex="female"
 /dev_stage="73 days post natal"
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ORIGIN

Query Match 18.8%; Score 167; DB 5; Length 535;
 Best Local Similarity 100.0%; Pred. No. 1e-78;
 Matches 167; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 721 ATGTGTGAGAACAACTGAGGAAAGACCTTCCCCACCCAGACGTTTTTATCTGA 780
 Db 535 ATGTGTGAGAACAACTGAGGAAAGACCTTCCCCACCCAGACGTTTTTATCTGA 476
 QY 781 ATGCATACAGGAGTCTCGAGTGGTGTGATTTGGCCAGTGTTAACCTGTGACAGTACT 840
 Db 475 ATGCATACAGGAGTCTCGAGTGGTGTGATTTGGCCAGTGTTAACCTGTGACAGTACT 416
 QY 841 CAGGTGTGAGGACAAAGTCAATGGCTCTCTCCCTGAGTGAAGAA 887
 Db 415 CAGGTGTGAGGACAAAGTCAATGGCTCTCTCCCTGAGTGAAGAA 369

RESULT 3

AI813346/c
 LOCUS AI813346 493 bp mRNA linear EST 21-DEC-1999
 DEFINITION wj33903.x1 NCI_CGAP_Kid12 Homo sapiens cDNA clone IMAGE:2404650 3',
 mRNA sequence.
 ACCESSION AI813346
 VERSION AI813346.1 GI:5424561
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 AUTHORS 1 (bases 1 to 493)
 TITLE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 JOURNAL National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 COMMENT Unpublished (1997)

Contact: Robert Strausberg, Ph.D.
 Email: csapbs@remail.nih.gov
 Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
 Emmert-Buck, M.D., Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D.
 cDNA Sequencing by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 www.bio.llnl.gov/bbrp/image/image.html
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 Seq primer: -40UP from Gibco
 High quality sequence stop: 447.
 Location/Qualifiers

FEATURES

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 /clone="IMAGE:2404650"
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 /note="Organ: kidney; Vector: pT7T3D-Pac (Pharmacia) with
 a modified polylinker; Site 1: Not 1; Site 2: Eco RI;
 Plasmid DNA from the normalized library NCI_CGAP kids was
 prepared, and ss circles were made in vitro. Following HAP
 purification, this DNA was used as tracer in a subtractive
 hybridization reaction. The driver was PCR-amplified cDNAs

/note="Organ: whole brain; Vector: Lafmid BA; Site 1: Not
 1; Site 2: Hind III; 1st strand cDNA was primed with a Not
 I - oligo(dT) primer [5',
 AACTGGAAGAAATTCGGCGCCGAGGAATTTTTTTTTTTT 3'];
 Double-stranded cDNA was ligated to Hind III adaptors
 (Pharmacia), digested with Not I and directionally cloned
 into the Not I and Hind III sites of the Lafmid BA vector.
 Library went through one round of normalization. Library
 constructed by Bento Soares and M.Fatima Bonaldo."


```

QY 383 CCGCTGGTACCCCGCGTGGAGAACTGGGCTTCAGCCGGGCTAAAGGA 435
Db 100 CCGCTGGTACCCCGCGTGGAGAACTGGGCTTCAGCCGGGCTAAAGGA 152

RESULT 6
R53538/c
LOCUS R53538 478 bp mRNA linear EST 18-MAY-1995
DEFINITION Y94c01.s1 Soares infant brain INIB Homo sapiens cDNA clones
ACCESSION R53538
VERSION IMAGE:40052 3', mRNA sequence.
KEYWORDS EST.
SOURCE R53538.1 GI:815440
ORGANISM Homo sapiens (human)

REFERENCE 1
AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaakis, E., Waterston, R., Williamson, A., Wohlmann, P. and Wilson, R.
TITLE The WashU-Merck EST Project
JOURNAL Unpublished (1995)
COMMENT 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Insert Size: 1472
High quality sequence stops: 402 Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1472 Std Error: 0.00
Seq primer: Promega -2imj3
High quality sequence stop: 402.

FEATURES
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1..478
Location/Qualifiers
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/db_xref="taxon:9606"
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/sex="female"
/dev_stage="73 days post natal"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares infant brain INIB"
/note="Organ: whole brain; Vector: Lafmid BA; Site 1: Not I; Site 2: Hind III; 1st strand cDNA was primed with a Not I - oligo(dT) primer [5' AACTGGAGAAATTCGGCGCGCAGGAATTTTTTTTTTTT 3']; double-stranded cDNA was ligated to Hind III adaptors (Pharmacia), digested with Not I and directionally cloned into the Not I and Hind III sites of the Lafmid BA vector. Library went through one round of normalization. Library constructed by Bento Soares and M.Fatima Bonaldo."

ORIGIN
Query Match 3.4%; Score 30; DB 7; Length 478;
Best Local Similarity 100.0%; Pred. No. 0.0024;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 832 ACAAGTACTCAGGTGTGAGACAAAGATGC 861
Db 431 ACAAGTACTCAGGTGTGAGACAAAGATGC 402

RESULT 7
CNS02COP/c
LOCUS CNS02COP 939 bp DNA linear GSS 01-SEP-2000

Query Match 3.4%; Score 30; DB 7; Length 478;
Best Local Similarity 100.0%; Pred. No. 0.0024;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 832 ACCCGCTTCTACCTGCCCAACTGTGAC 706
Db 503 ACCCGCTTCTACCTGCCCAACTGTGAC 477

RESULT 8
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LOCUS R85550 316 bp mRNA linear EST 14-AUG-1995
DEFINITION Y038d07.s1 Soares adult brain N2b4HB55Y Homo sapiens cDNA clone
ACCESSION R85550
VERSION IMAGE:180205 3', mRNA sequence.
KEYWORDS EST.
SOURCE R85550.1 GI:943956
ORGANISM Homo sapiens (human)

REFERENCE 1
AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M.,

Tetraodon nigroviridis genome survey sequence T7 end of clone
254D23 of library G from Tetraodon nigroviridis, genomic survey
sequence.
AL190546
GI:7828650
GSS; genome survey sequence.
Tetraodon nigroviridis
Tetraodon nigroviridis
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Acanthopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
Acanthopterygii; Acanthopterygii; Percomorpha; Tetraodontiformes;
Tetraodontidae; Tetraodontidae; Tetraodon.
1
Roest Crolius, H., Jaillon, O., Dasilva, C., Bouneau, L., Fisher, C.,
Bernot, A., Fzanes, C., Wincker, P., Brotier, P., Quetier, F.,
Saurin, W. and Weissenbach, J.
Estimate of human gene number provided by genome-wide analysis
using Tetraodon nigroviridis DNA sequence
Nat. Genet. 25 (2), 235-238 (2000)
20296633
10835645
2
Roest Crolius, H., Jaillon, O., Dasilva, C., Ozouf-Costaz, C.,
Fizames, C., Fischer, C., Bouneau, L., Billault, A., Quetier, F.,
Saurin, W., Bernot, A. and Weissenbach, J.
Characterization and repeat analysis of the compact genome of the
freshwater pufferfish Tetraodon nigroviridis
Genome Res. 10 (7), 939-949 (2000)
20359837
10899143
3 (bases 1 to 939)
Genoscope.
Direct Submission
Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :
BP 191 91006 Evry cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)
- Web : www.genoscope.cns.fr)
This sequence is a single read and was generated as part of a large
scale clone-end sequencing project of the tetraodon nigroviridis
genome. For more information, please take a look at
http://www.genoscope.cns.fr/Tetraodon.
Location/Qualifiers
1..939
/organism="Tetraodon nigroviridis"
/mol_type="Genomic DNA"
/db_xref="taxon:99883"
/clone="254D23"
/clone_lib="G"
/note="Genoscope sequence ID : COAG254CB12LP1-end : T7"

Query Match 3.0%; Score 27; DB 9; Length 939;
Best Local Similarity 100.0%; Pred. No. 0.01;
Matches 27; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 680 ACCCGCTTCTACCTGCCCAACTGTGAC 706
Db 503 ACCCGCTTCTACCTGCCCAACTGTGAC 477

RESULT 8
R85550
LOCUS R85550 316 bp mRNA linear EST 14-AUG-1995
DEFINITION Y038d07.s1 Soares adult brain N2b4HB55Y Homo sapiens cDNA clone
ACCESSION R85550
VERSION IMAGE:180205 3', mRNA sequence.
KEYWORDS EST.
SOURCE R85550.1 GI:943956
ORGANISM Homo sapiens (human)

REFERENCE 1
AUTHORS Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M.,

```

TITLE
JOURNAL
COMMENT

Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M.,
 Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F.,
 Trevastis, E., Waterston, R., Williamson, A., Wohldmann, P. and
 Wilson, R. Merck EST Project
 Unpublished (1995)
 Contact: Wilson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 Insert Size: 1578
 High quality sequence stops: 278
 Source: IMAGE Consortium, LLNL
 This clone is available royalty-free through LLNL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Insert length: 1578 Std Error: 0.00
 Seq primer: Promega -21ml3
 High quality sequence stop: 278.
 Location/Qualifiers
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 /mol_type="mRNA"
 /db_xref="GBB:3826849"
 /db_xref="taxon:9606"
 /clone="IMAGE:180205"
 /sex="Male"
 /dev_stage="55-Year old"
 /lab_host="DH10B (ampicillin resistant)"
 /clone_lib="Soares adult brain N2b4HB55y"
 /note="Organ: brain; Vector: pTT73D (Pharmacia) with a
 modified polylinker; Site 1: Not I; Site 2: Eco RI; 1st
 strand cDNA was primed with a Not I - oligo(dT) primer [5'
 TGTTCACATCTGAAGTGGAGCGCGCTTTTCTTTTCTTTT 3'],
 double-stranded cDNA was size selected, ligated to Eco RI
 adapters (Pharmacia), digested with Not I and cloned into
 the Not I and Eco RI sites of a modified pTV73 vector
 (Pharmacia). Library went through one round of
 normalization to a Cot = 53. Library constructed by Bento
 Soares and M. Fatima Bonaudo. The adult brain RNA was
 provided by Dr. Donald H. Gilden. Tissue was acquired
 17-18 hours after death which occurred in consequence of a
 ruptured aortic aneurysm. RNA was prepared from a pool of
 tissues representing the following areas of the brain:
 frontal, parietal, temporal and occipital cortex from the
 left and right hemispheres, subcortical white matter,
 basal ganglia, thalamus, cerebellum, midbrain, pons and
 medulla."

FEATURES
 source

1..601
 /organism="Bos taurus"
 /mol_type="genomic DNA"
 /strain="breed: Hereford"
 /db_xref="taxon:9913"
 /clone="CH240_211K10"
 /sex="Male"
 /cell_type="Blood"
 /clone_lib="CHORI-240"
 /note="Vector: pTARBAC1.3; Site 1: MboI; Site 2: MboI;
 Hereford bull L1 Domino 99375; CHORI-240 Bovine BAC
 library (Male) produced by Pieter de Jong"

ORIGIN

Query Match 2.5%; Score 22; DB 8; Length 601;
 Best Local Similarity 100.0%; Pred. No. 5.3; Indels 0; Gaps 0;
 Matches 22; Conservative 0; Mismatches 0;

QY 744 GAAAGCACCTTCCCCCACCACA 765
 DB 229 GAAAGCACCTTCCCCCACCACA 250

RESULT 10
AZ617122/c
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

AZ617122 441 bp DNA linear GSS 13-DEC-2000
 1M0448D17F Mouse 10kb plasmid UUGC1M library Mus musculus genomic
 clone UUGC1M0448D17 F, genomic survey sequence.

AZ617122 GI:11739312
 GSS.
 Mus musculus (house mouse)

REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 1 (bases 1 to 441)
 Dunn, D., Aoyagi, A., Barber, M., Beacorn, T., Duval, B., Hamil, C.,
 Islam, H., Longacre, S., Mahmoud, M., Meenen, E., Pedersen, I.,
 Reilly, M., Rose, R., Rose, R., Stokes, R., Tingey, A., von
 Niederhausern, A. and Wright, D., Weiss, R.
 Mouse whole genome scaffolding with paired end reads from 10kb
 plasmid inserts
 Unpublished (2000)
 Contact: Robert B. Weiss
 University of Utah Genome Center
 University of Utah
 Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLC, UT
 84112, USA

QY 512 GCCTGGCTGACGGCTCAGGGCG 533
 DB 212 GCCTGGCTGACGGCTCAGGGCG 233

RESULT 9
BZ869242
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

BZ869242 601 bp DNA linear GSS 18-MAR-2003
 CH240_211K10_TJ CHORI-240 Bos taurus genomic clone CH240_211K10,
 genomic survey sequence.
 BZ869242 GI:29096647
 GSS.
 Bos taurus (cow)
 Bos taurus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 Bovinae; Bos.

Tel: 801 585 5606
 Fax: 801 585 7177
 Email: dgunn@genetics.utah.edu
 Insert Length: 10000 Std Error: 0.00
 Plate: 0448 row: D column: 17
 Seq primer: CGTGTAAACGACGGCCAGT
 Class: plasmid ends
 High quality sequence stop: 441.

FEATURES

Location/Qualifiers
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 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="UUGC1M0448D1.7"
 /sex="Male"

/lab_host="E. Coli strain XL10-Gold, TI-resistant, F-"
 /clone_lib="Mouse 10kb plasmid UUGC1M library"
 /notes="vector: PWD42nv; Purified genomic DNA from M.
 musculus C57BL/6J (male) was obtained from the Jackson
 Laboratory Mouse DNA Resource
 (http://www.jax.org/resources/documents/dnares/). The DNA
 was hydrodynamically sheared by repeated passage through a
 0.005 inch orifice at constant velocity. The sheared DNA
 was blunt end-repaired with T4 DNA polymerase and T4
 polynucleotide kinase. Adaptor oligonucleotides were
 ligated to the blunt ends in high molar excess. The
 adaptor DNA was purified and size-selected for a 9.5 to
 10.5 kb range using preparative agarose gel
 electrophoresis. Vector DNA was prepared from a derivative
 of pW42 (G14732114|GB|AF129072.1), a copy-number
 inducible derivative of plasmid R1. The vector was ligated
 with adaptors complementary to the insert adaptors and
 purified. The sheared, adaptor mouse DNA was annealed to
 adaptor vector DNA, and transformed into
 chemically-competent E. coli XL10-Gold (Stratagene) cells
 and selected for ampicillin resistance."

ORIGIN

Query Match 2.4%; Score 21; DB 8; Length 441;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 586 GAGAAACACCTGTCAGGC 606

Db 276 GAGAAACACCTGTCAGGC 256

RESULT 11

BY590905/c 462 bp mRNA linear EST 15-DEC-2002
 LOCUS BY590905 RIKEN full-length enriched, adult inner ear Mus musculus
 DEFINITION cDNA clone F930014102 3', mRNA sequence.

ACCESSION BY590905

VERSION BY590905.1

KEYWORDS EST.

SOURCE Mus musculus (house mouse)

ORGANISM Mus musculus

REFERENCE 1 (bases 1 to 462)
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS

Nikaido, I., Furuno, M., Kasukawa, T., Adachi, J., Bono, H., Kondo, S.,
 Kiyosawa, H., Yagi, K., Tomaru, Y., Hasegawa, Y., Nogami, A.,
 Schonbach, C., Gojobori, T., Baldarelli, R., Hill, D. P., Buit, C.,
 Hume, D. A., Quackenbush, J., Schriml, L. M., Kanapin, A., Matsuda, H.,
 Batalov, S., Beisel, K. W., Blake, J. A., Brad, D., Brusci, V.,
 Chothia, C. P., Cousani, L. E., Cousins, S., Dalla, E., Dragani, T. A.,
 Fletcher, C. P., Forrest, A., Frazer, K. S., Gaasterland, T.,
 Gariboldi, M., Gissi, C., Godzik, A., Gough, J., Grimmond, S.,
 Gustincich, S., Hirokawa, N., Jackson, I. J., Jarvis, E. D., Kanai, A.,
 Kawaji, H., Kawasawa, Y., Kedierski, R. N., King, B. L., Koragaya, A.,
 Kurochkin, I. V., Lee, Y., Lenhard, B., Lyons, P. A., Maglott, D. R.,

Maltais, L., Marchionni, L., McKenzie, L., Miki, H., Nagashima, T.,
 Numata, K., Okido, T., Pavan, W. J., Perte, G., Pesole, G.,
 Petrovsky, N., Pillai, R., Pontius, J. U., Qi, D., Ramchandran, S.,
 Ravasi, T., Reed, J. C., Reed, D. J., Reid, J., Ring, B. Z., Ringwald, M.,
 Sandelin, A., Schneider, C., Semple, C. A., Setou, M., Shimada, K.,
 Sultana, R., Takemura, Y., Taylor, M. S., Teasdale, R. D., Tomita, M.,
 Verardo, R., Wagner, L., Wahlestedt, C., Wang, Y., Watanabe, Y.,
 Wells, C., Wilming, L. G., Wynshaw-Boris, A., Yanagisawa, M., Yang, I.,
 Yang, L., Yuan, Z., Zavolan, M., Zhu, Y., Zimmer, A., Carninci, P.,
 Hayatsu, N., Hirozane-Kishikawa, T., Konno, H., Nakamura, M.,
 Sakazume, N., Sato, K., Shiraki, T., Waki, K., Kawai, J., Azawa, K.,
 Arakawa, T., Fukuda, S., Hara, A., Hashizume, W., Inotani, K., Ishii, Y.,
 Itch, M., Kagawa, I., Miyazaki, A., Sakai, K., Sasaki, D., Shibata, K.,
 Shinagawa, A., Yasunishi, A., Yoshino, M., Waterston, R., Lander, E. S.,
 Rogers, J., Birney, E. and Hayashizaki, Y.

TITLE

Analysis of the mouse transcriptome based on functional annotation
 of 60,770 full-length cDNAs

JOURNAL

Nature 420, 563-573 (2002)

MEDLINE

22354683

PUBMED

12466851

COMMENT

Contact: Yoshihide Hayashizaki
 Laboratory for Genome Exploration Research Group, RIKEN Genomic
 Sciences Center (GSC), Yokohama Institute
 The Institute of Physical and Chemical Research (RIKEN)
 1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
 Tel: 81-45-503-9222
 Fax: 81-45-503-9216

Email: genome-res@gsc.riken.jp, URL: http://genome.gsc.riken.jp/
 Aizawa, K., Akimura, T., Arakawa, T., Carninci, P., Fukuda, S.,
 Hirozane, T., Inotani, K., Ishii, Y., Itoh, M., Kawai, J., Konno, H.,
 Miyazaki, A., Murata, M., Nakamura, M., Nomura, K., Numazaki, R.,
 Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K.,
 Shiraki, T., Tagami, M., Waki, K., Watahiki, A., Muramatsu, M. and
 Hayashizaki, Y. Direct Submission

Computational Analysis of Full-Length Mouse cDNAs Compared with
 Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
 Normalization and subtraction of cap-trapper-selected cDNAs to
 prepare full-length cDNA libraries for rapid discovery of new
 genes. Genome Res. 10 (10), 1617-1630 (2000)
 RIKEN integrated sequence analysis (RISA) system--384-format
 sequencing pipeline with 384 multicapillary sequencer. Genome Res.
 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA
 encyclopedia: real-time sequence clustering for construction of a
 nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
 cDNA library was prepared and sequenced in Mouse Genome
 Encyclopedia Project of Genome Exploration Research Group in Riken
 Genomic Sciences Center and Genome Science Laboratory in RIKEN,
 Division of Experimental Animal Research in Riken contributed to
 Prepare mouse tissues.
 Tissues were provided by Kirk W. Beisel (Boys Town National
 Research Hospital 555 North 30th Street Omaha, NE 68131 USA) whose
 assistance we gratefully acknowledge.
 Please visit our web site (http://genome.gsc.riken.go.jp) for
 further details.

FEATURES

Location/Qualifiers
 1..462
 /organism="Mus musculus"
 /mol_type="mRNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="F930014102"
 /tissue_type="inner ear"
 /dev_stage="adult"
 /clone_lib="RIKEN full-length enriched, adult inner ear"

ORIGIN

Query Match 2.4%; Score 21; DB 5; Length 462;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 748 GCACCTTCCCCCAGCCAGAC 768

Db 262 GCACCTTCCCCCAGAC 242

RESULT 12
AQ294438
LOCUS
DEFINITION HS 3014 B2.H04 T7 CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=3014 Col=8 Row=P, genomic survey sequence.
ACCESSION AQ294438.1 GI:4012021
VERSION AQ294438
KEYWORDS GSS.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 479)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D., and Hood,L.

TITLE Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
MEDLINE 93180589
PUBMED 1049764

COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3014 row: P column: 8
Class: PAC ends
High quality sequence stop: 479.

FEATURES
source Location/Qualifiers
1..479
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/clone="Plate=3014 Col=8 Row=P"
/sex="male"
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/note="Organ: sperm; Vector: pBeloBAC11; PAC Clones in E-Coli DH10B"

ORIGIN
Query Match 2.4%; Score 21; DB 8; Length 479;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 745 AAAGCACCTTCCCCCAGAC 765
Db 219 AAAGCACCTTCCCCCAGAC 239

RESULT 13
AA823641
LOCUS
DEFINITION AA823641 494 bp mRNA linear EST 17-FEB-1998
IMAGE:1125882 5', mRNA sequence.
ACCESSION AA823641
VERSION AA823641.1 GI:2893509
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 494)
Marra,M., Hallier,L., Allen,M., Bowles,M., Dietrich,N., Dubuque,T., Geisler,S., Kucaba,T., Lacy,M., Le,M., Martin,J., Morris,M., Schellenberg,K., Seftoe,K., Tan,F., Underwood,K., Moore,B.,

Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and Waterston,R.
The WashU-HMI Mouse EST Project
Unpublished (1996)
Contact: Marra M/Mouse EST Project
WashU-HMI Mouse EST Project
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.wustl.edu
This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.
MG1:615218
High quality sequence stop: 493.

FEATURES
source Location/Qualifiers
1..494
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/mol_type="mRNA"
/strain="C57BL/6J x DBA/2J F1"
/db_xref="taxon:10090"
/clone="IMAGE:1125882"
/tissue_type="embryo"
/dev_stage="2-cell"
/lab_host="DH10B"
/clone_lib="Knowles Solter mouse 2 cell"
/note="Organ: embryo; Vector: pBluescribe (modified); Site 1: Mui; Site 2: Sali; Cloned unidirectionally from mRNA prepared from 13,500 2-cell stage embryos. Primer: Sali (dr): 5'-CGTGCACCGTCGACCGTTTCTTTT-3'. CDNAS were cloned into the Mui/Sali sites of a modified pBluescribe vector using commercial linkers (NEB). Average insert size: 1.2 kb."

ORIGIN
Query Match 2.4%; Score 21; DB 1; Length 494;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 748 GCACCTTCCCCCAGAC 768
Db 365 GCACCTTCCCCCAGAC 385

RESULT 14
BB628603
LOCUS
DEFINITION BB628603 657 bp mRNA linear EST 26-OCT-2001
musculus cDNA clone 9630018N03 5', mRNA sequence.
ACCESSION BB628603
VERSION BB628603.1 GI:16465993
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 657)
Hara,A., Hiramoto,K., Hori,F., Ishii,Y., Ito,M., Kawai,J., Konno,H., Kouda,M., Koya,S., Matsuyama,T., Miyazaki,A., Nomura,K., Ohno,M., Okazaki,Y., Okido,T., Saito,R., Sakai,C., Sakai,K., Sano,H., Sasaki,D., Shibata,K., Shinagawa,A., Shiraki,T., Sogabe,Y., Suzuki,H., Tagami,M., Tagawa,A., Takahashi,F., Takeda,Y., Tanaka,T., Toya,T., Muramatsu,M. and Hayashizaki,Y.
RIKEN Mouse ESTs (Arakawa,T., et al. 2001)
Unpublished (2001)

TITLE Yoshihide Hayashizaki
JOURNAL Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
COMMENT The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-resseqc.riken.jp, URL: <http://genome.gsc.riken.jp/>
 Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh, M., Kono, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
 Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. *Genome Res.* 10 (10), 1617-1630 (2000)
 wagi, K., Fujiwara, S., Inoue, K., Togawa, Y., Izawa, M., Chara, E., Watabiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura, S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and Hayashizaki, Y.
 RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. *Genome Res.* 10 (11), 1757-1771 (2000)
 Kono, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara, Y. and Hayashizaki, Y.
 Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. *Genome Res.* 11 (2), 281-289 (2001)
 Kondo, S., Shinagawa, A., Saito, T., Kiyosawa, H., Yamakawa, T., Aizawa, K., Fukuda, S., Hara, A., Itoh, M., Kawai, J., Shibata, K. and Hayashizaki, Y.
 Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences. *Mamm. Genome* 12, 673-677 (2001)
 Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES

source
 1. .657
 /organism="Mus musculus"
 /mol_type="mRNA"
 /db_xref="taxon:10090"
 /clone="9630018N03"
 /tissue_type="cerebellum"
 /dev_stage="16 days neonate"
 /lab_host="DH10B"
 /clone_lib="RIKEN full-length enriched, 16 days neonate cerebellum"
 /note="Site 1: SalI; Site 2: BamHI; cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues. 1st strand cDNA was primed with a primer [5', GAGGAGAGAGATCCAGAGCTTTTCTTTTCTTTTNN 3'], cDNA was prepared by using trehalose thermo-activated reverse transcriptase and subsequently enriched for full-length by cap-trapper. cDNA went through one round of normalization to Rot = 20.0 and subtraction to Rot = 370.0. Second strand cDNA was prepared with the primer adapter of sequence [5', GAGGAGAGATCTCCAGTTAATAAATTAATCCCCCCCC 3']. cDNA was cleaved with XhoI and BamHI. vector: a modified pBluescript KS(+) after bulk excision from Lambda FLC I."

ORIGIN

Query Match 2.4%; Score 21; DB 2; Length 657;
 Best Local Similarity 100.0%; Pred. No. 18;
 Matches 21; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 748 GCACCTTCCCCCCCCCAGAC 768
 Db 75 GCACCTTCCCCCCCCCAGAC 95

RESULT 15

CNS0413S/c
 LOCUS 803 bp DNA linear GSS 01-SEP-2000
 DEFINITION Tetraodon nigroviridis genome survey sequence PUC-Ori end of clone 073p11 of library G from Tetraodon nigroviridis, genomic survey sequence.
 ACCESSION AL269713
 VERSION AL269713.1 GI:7991606

KEYWORDS

GSS; genome survey sequence.

SOURCE

Tetraodon nigroviridis
 Tetraodon nigroviridis
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Acanthopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Acanthopterygia; Acanthopterygii; Percomorpha; Tetraodontiformes; Tetraodontidae; Tetraodontidae; Tetraodon.

REFERENCE

1
 Roest Crolius, H., Jaillon, O., Dasilva, C., Bouneau, L., Fisher, C., Bernot, A., Pizanes, C., Wincker, P., Brottier, P., Quetier, F., Saurin, W., and Weissenbach, J
 Estimate of human gene number provided by genome-wide analysis using Tetraodon nigroviridis DNA sequence
Nat. Genet. 25 (2), 235-238 (2000)

JOURNAL

MEDLINE
 20296633

PUBMED

10835645

AUTHORS

2
 Roest Crolius, H., Jaillon, O., Dasilva, C., Ozouf-Costaz, C., Pizanes, C., Fischer, C., Bouneau, L., Billault, A., Quetier, F., Saurin, W., Bernot, A., and Weissenbach, J.
 Characterization and repeat analysis of the compact genome of the freshwater pufferfish Tetraodon nigroviridis
Genome Res. 10 (7), 939-949 (2000)

JOURNAL

MEDLINE
 20359837

PUBMED

10899143

AUTHORS

3 (Bases 1 to 803)

TITLE

Direct Submission

JOURNAL

Submitted (12-APR-2000) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : secref@genoscope.cns.fr)
 - Web : www.genoscope.cns.fr
 This sequence is a single read and was generated as part of a large scale clone-end sequencing project of the Tetraodon nigroviridis genome. For more information, please take a look at <http://www.genoscope.cns.fr/Tetraodon>.

FEATURES

source

1. .803
 /organism="Tetraodon nigroviridis"

/mol_type="genomic DNA"

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ORIGIN

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 Db 774 CTAATGACAGATGTGTGAGA 754

Search completed: November 7, 2004, 06:03:22

Job time : 4554 secs